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818-1320 of SEQ ID NO:3

368--1144 of SEQ ID NO:34

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Searcher: _____
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Searcher Prep Time: _____
Online Time: _____

Type of Search
NA# 3 AA#: _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure #: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable
STN: _____
DIALOG: _____
QUESTEL/ORBIT: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: 04
WWW/Internet: _____
Other (Specify): _____

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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:52:53 ; Search time 2639.94 Seconds
(without alignments)
9232.381 Million cell updates/sec

Title: US-09-463-542-3_COPY_818_1320

Perfect score: 503

Sequence: 1 tagagtaagacaccttggaa.....atatattggaactgatgtct 503

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_hcg.*
3: gb_in.*
4: gb_on.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pi.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	473	94.0	597	9	AF012874 Homo sapi
2	455.2	90.5	984	9	AF310249 Homo sapi
3	455.2	90.5	1100	6	AF121474 Sequence
4	455.2	90.5	1100	9	AB005520 Homo sapi
5	455.2	90.5	85873	6	AX951686 Sequence
6	455.2	90.5	135675	9	AC093174 Homo sapi
7	455.2	90.5	148828	9	AY157024 Homo sapi
8	455.2	90.5	166049	9	AC090947 Homo sapi
9	362.8	72.1	186028	2	AC151849 Callithr
10	219.6	43.7	191540	2	AC136055 Rattus no
11	219.6	43.7	245724	2	AC091418 Rattus no
12	219.6	43.7	250169	2	AC120668 Rattus no
13	216.4	43.0	682	10	S79407 mPPAR gamma
14	216.4	43.0	1159	10	AY243584 Mus muscu
15	216.4	43.0	1160	10	AY236531 Mus muscu
16	216.4	43.0	185737	2	AC125447 Mus muscu
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31	44.4	8.8	165260	9	AC024341
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ALIGNMENTS

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DEFINITION	AF012874	Homo sapiens	PPARG gene, isoform 2,	promoter and 5'UTR	sequence.	
ACCESSION	AF012874	Homo sapiens	PPARG gene, isoform 2,	promoter and 5'UTR	sequence.	
VERSION	AF012874.1	GI:4731428				
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
MEDLINE						
PUBMED						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
FEATURES						
source						
gene						
promoter						
5'UTR						
ORIGIN						

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Best Local Similarity 100.0%; Pred. No. 5e-101;
Matches 473; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 151 TGCCCAATAAGCTTTCTGTGATTTTATATGATGATGATGATGATGATGATGATGATGATGAT 210
Db 121 TGCCCAATAAGCTTTCTGTGATTTTATATGATGATGATGATGATGATGATGATGATGATGAT 180

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Db 181 CCATTGTGAATTTATACAAACAATAAATAAATGCAAGTGGATATTTGAACAGTCTCTCTCTGA 240

Qy 271 TAAATCTTAAATACAGTACAGTTCACGCCCTCAGGACACCTGACATGTTGGTCAACCGGC 330
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Qy 331 GAGACAGTGTGGCAATATTTATCCCTGTAATGTACCAAGTCTTGGCAGACAGTGAACATT 390
Db 301 GAGACAGTGTGGCAATATTTATCCCTGTAATGTACCAAGTCTTGGCAGACAGTGAACATT 360

Qy 391 ATGACACAACCTTTTGTGTCAGAGTGGCTCTCTTAATAGGACAGTGCAGGCAATTTCAAGCCC 450
Db 361 ATGACACAACCTTTTGTGTCAGAGTGGCTCTCTTAATAGGACAGTGCAGGCAATTTCAAGCCC 420

Qy 451 AGTCCCTTCTGTGTTTATTTCCCATCTCTCCCAATATTTTGAACAGTGAATGCT 503
Db 421 AGTCCCTTCTGTGTTTATTTCCCATCTCTCCCAATATTTTGAACAGTGAATGCT 473

RESULT 2
AF310249
LOCUS Homo sapiens peroxisome proliferator activated receptor gamma 2
DEFINITION gene, upstream sequence.
ACCESSION AF310249
VERSION AF310249.1 GI:13274397
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE Lim, C.H., Kok, C.C., Samian, M.R., Najimudin, N. and Tengku
Muhammad, T.S.
Molecular cloning and sequencing of the human Peroxisome
Proliferator Activated Receptor Gamma 2 promoter
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 984)
AUTHORS Lim, C.H., Kok, C.C., Samian, M.R., Najimudin, N. and Tengku
Muhammad, T.S.
Direct Submission
JOURNAL Submitted (03-OCT-2000) School of Biological Sciences, Universiti
Sains Malaysia, Minden, Penang 11800, Malaysia
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source 1..984
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2"
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Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

Qy 1 TAGAGTAGTAGTACCTTAGGAATATACCAATTTTCAGTAGCATGCTGATACCAAGCTTTAAACT 60
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Qy 61 ATGATATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGTGATTTAGAGATTAGAGATT 120
Db 422 ATGATATACATATTTGAATTTCCAAATTTTCTTCAATAATGT-----GATTAGAGATT 474

Qy 121 CAACCCAGGGATAGACACCGGAAGAAACTTTTGGCCCAATTAAGCTTTCTGGTATTTTCATAA 180
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Qy 181 GCAAGAGATTTAAGTTTTCCATTTTAAGAAGCAATTTGGCCCAATTAAGCTTTCTGGTATTTTCATAA 240
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Qy 301 TCACGACAGACTGNAACATGT-GGTCACCGGCGGACAGTGTGGCAATATTATCCCTGTAA 359
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Qy 360 TGTACCAAGTCTTGGCCAGAGCAGTGAACATTTATGACACAACTTTTGTTCACAGCTGGCTC 419
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Qy 480 CCAAATATTTGGAACTGATGCTCT 503
Db 835 CCAAATATTTGGAACTGATGCTCT 858

RESULT 3
AR121474
LOCUS Sequence 10 from patent US 6159734.
DEFINITION AR121474
ACCESSION AR121474
VERSION AR121474.1 GI:14105050
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 1100)
AUTHORS McKay, R., Borchers, A.H. and Baker, B.F.
TITLE Antisense modulation of peroxisome proliferator-activated receptor
gamma expression
JOURNAL Patent: US 6159734-A 10 12-DEC-2000;
FEATURES Location/Qualifiers
source 1..1100
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Query Match      90.5%; Score 455.2; DB 6; Length 1100;
Best Local Similarity 96.8%; Pred. No. 7.3e-97;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;
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Qy 1 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAAGCTTTAAACT 60
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Db 352 CAACAGGAGATAGACACCGAAGAAACTTTGCCCAATAAGCTTTCTGCTATTTTCATAA 411
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Qy 360 TGTACCAAGCTTTGCCAGAGCAGTGAACATTTATGACACAACTTTTGTGACAGCTGGCTC 419
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Db 712 CCAAATATTTGGAAACTGATGCT 735

RESULT 4
LOCUS AB005520 1100 bp DNA linear PRI 10-NOV-1997
DEFINITION Homo sapiens ppar gamma2 gene for peroxisome proliferator
activated-receptor gamma, partial cds and 5' flanking.
ACCESSION AB005520
VERSION AB005520.1 GI:2605488
KEYWORDS ppar gamma2; peroxisome proliferator activated-receptor gamma.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
AUTHORS Okazawa,H., Mori,H., Tamori,Y., Araki,S., Niki,T., Masugi,J.,
Kawanishi,M., Kubota,T., Sinoda,H. and Kasuga,M.
TITLE No coding mutations are detected in the peroxisome
proliferator-activated receptor- gene in Japanese patients with
lipotrophic diabetes
JOURNAL Diabetes (1997) In press
REFERENCE 2 (bases 1 to 1100)
AUTHORS Okazawa,H.
TITLE Direct Submission
JOURNAL Submitted (03-JUL-1997) Hideki Okazawa, Kobe University School of
Medicine, 2nd Department of Internal Medicine; 7-5-1 Kusunoki-cho
Chuo-ku, Kobe 650, Japan (E-mail:okazawa@med.kobe-u.ac.jp,
Tel:81-78-341-7451, Fax:81-78-382-2080)
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Query Match 90.5%; Score 455.2; DB 9; Length 1100;
Best Local Similarity 96.8%; Pred. No. 7.3e-97;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;
Qy 1 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAAGCTTTAAACT 60
Db 239 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAAGCTTTAAACT 298
Qy 61 ATGATACATATTTGAATTTCCAAATTTTCTTCAGTAATGCTGATAGAGATTAGAGATT 120
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Qy 181 GCAAGAGATTTAAGTTTTCCAAATTTTGAAGCCATTTGTAATATTAACAAATAAATG 240
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Qy 241 CAAGTGGATATTAACAGCTCTCTCTCTGATAATTTCTAAATACAGTACAGTTCACGCC 300
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Qy 480 CCAAATATTTGGAAACTGATGCT 503
Db 712 CCAAATATTTGGAAACTGATGCT 735

RESULT 5
LOCUS AX951686/c 85873 bp DNA linear PAT 09-JAN-2004
DEFINITION Sequence 3 from Patent WO03093310.
ACCESSION AX951686
VERSION AX951686.1 GI:40782040
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE
AUTHORS Bullerdick,J.
TITLE Novel nucleic acid sequences and proteins of tumors and neoplasias
of the thyroid gland
JOURNAL Patent: WO 03093310-A 3 13-NOV-2003;
Universitaet Bremen (DE)
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Qy	61	ATGGATACATATTTGAAATTCCAAATTTTCTCTCAGATAATCTGATTAGAGATTAGAGATT	120
Db	54412	ATGGATACATATTTGAAATTCCAAATTTTCTCTCAGATAATCTGATTAGAGATTAGAGATT	54360
Qy	121	CAACAGGATAGACACCGAAGAAACTTTTGGCCAAATAAGCTTTCTGGTATTTCATAA	180
Db	54359	CAACAGGATAGACACCGAAGAAACTTTTGGCCAAATAAGCTTTCTGGTATTTCATAA	54300
Qy	181	GCAAGAGATTTAAGCTTTTCCATTTAAGAGCCATTGTGAATTTATACACATTAATAATG	240
Db	54299	GCAAGAGATTTAAGCTTTTCCATTTAAGAGCCATTGTGAATTTTACACATTAATAATG	54240
Qy	241	CAAGTGGATATTGAACAGTCTCTCTCTGATAATTTCTAAATACAGTTCACGCCCC	300
Db	54239	CAAGTGGATATTGAACAGTCTCTCTCTGATAATTTCTAAATACAGTTCACGCCCC	54180
Qy	301	TCACAGACATGAAACATGTT-GGTACCGCGGAGACAGTGTGGCAATATTATCCCTGTAA	359
Db	54179	TCACAAGACATGAAACATGTTGGGTACCGCGGAGACAGTGTGGCAATATTATTCCTCTGAA	54120
Qy	360	TGTACCAAGTCTGCCAGAGCAGTGAACATTTATGACACAACTTTTTCAGCTGGCTC	419
Db	54119	TGTACCAAGTCTTGCCTAAAGCAGTGAACATTTATGACACAACTTTTTCAGCTGGCTC	54060
Qy	420	CTAATAGACAGTGCACGCCAATTCAGCCAGTCCCTTTCTGTGTTATTCCTCATCTCTC	479
Db	54059	CTAATAGACAGTGCACGCCAATTCAGCCAGTCCCTTTCTGTGTTATTCCTCATCTCTC	54000
Qy	480	CCAAATATTGGAACTGATGTCT 503	
Db	53999	CCAAATATTGGAACTGATGTCT 53976	
RESULT 6		AC093174 135675 bp DNA linear PRI 08-NOV-2002	
AC093174		Homo sapiens chromosome 3 clone RP11-167M22 map 3p, complete	
LOCUS		sequence.	
DEFINITION		AC093174.2 GI:24796717	
ACCESSION		HTG.	
VERSION		Homo sapiens (human)	
KEYWORDS		Homo sapiens	
SOURCE		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
ORGANISM		Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
REFERENCE		1 (bases 1 to 135675)	
AUTHORS		Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J., Ding, H., Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D., Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C., Li, F., Li, G., Li, J., Li, L., Li, S., Li, T., Liu, Y., Liu, N., Liu, B., Liu, Y., Li, W., Li, W., Li, Y., Luo, J., Niu, Y., Qi, O., Qi, X., Song, L., Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J., Wang, J., Wang, L., Wang, L., Wang, R., Wang, R., Wang, X., Wang, X., Wang, Y., Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B., Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M., Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N., Yu, J. and Yang, H.	
TITLE		Chromosome 3p genomic sequence	
JOURNAL		Unpublished	
REFERENCE		2 (bases 1 to 135675)	
AUTHORS		Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J., Ding, H.,	
TITLE		Submitted (13-AUG-2001) Human Genomic Center, Institute of	
JOURNAL		Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing	
REFERENCE		100101, P.R.China	
AUTHORS		3 (bases 1 to 135675)	
TITLE		Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J., Ding, H., Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D., Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C., Li, F., Li, G., Li, J., Li, L., Li, S., Li, T., Liu, Y., Liu, N., Liu, B., Liu, Y., Li, W., Li, W., Li, Y., Luo, J., Niu, Y., Qi, O., Qi, X., Song, L., Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J., Wang, J., Wang, L., Wang, L., Wang, R., Wang, R., Wang, X., Wang, X., Wang, Y., Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B., Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M., Zhang, X., Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N., Yu, J. and Yang, H.	
TITLE		Direct Submission	
JOURNAL		Submitted (08-NOV-2002) Human Genomic Center, Institute of	
COMMENT		Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing	
		100101, P.R.China	
		On Nov 8, 2002 this sequence version replaced gi:15148930.	
		-----Genome Center	
		Center:Beijing Center	
		Center code:Beijing	
		Website:http://hgci.gtp.ac.cn	
		http://www.genomics.org.cn	
		Contact:hgci.gtp.ac.cn	
		----- Project Information	
		Center project name:1k project	
		Center clone name: RP11-167M22	
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		Consensus quality: 0 bases at least Q30	
		Consensus quality: 6 bases at least Q20	
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FEATURES	source	Location/Qualifiers	
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		/map="3p"	
		/clone="RP11-167M22"	
ORIGIN	Query Match	90.5%; Score 455.2; DB 9; Length 135675;	
	Best Local Similarity	96.8%; Pred. No. 4.5e-97;	
	Matches 488; Conservative	0; Mismatches 8; Indels 8; Gaps 2;	
Qy	1	TAGAGTAGTACCTTAGGAATATACATTTTCAGTAGCAGTGTGATACCAAGTTTAACT	60
Db	81198	TAGAGTAGTACCTTAGGAATATACATTTTCAGTAGCAGTGTGATACCAAGTTTAACT	81257
Qy	61	ATGGATACATATTTGAAATTCCAAATTTTCTCTCAGATAATCTGATTAGAGATTAGAGATT	120
Db	81258	ATGGATACATATTTGAAATTCCAAATTTTCTCTCAGATAATCTGATTAGAGATTAGAGATT	81310


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    /replace="c"
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    /rpt_type=dispersed
repeat_region
8735. .9019
    /rpt_family="Alu"
    /rpt_type=dispersed
repeat_region
9637. .9935
    /rpt_family="Alu"
    /rpt_type=dispersed
repeat_region
10092. .10404
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repeat_region
10423. .10537
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repeat_region
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repeat_region
22582. .22916
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Query Match      90.5%; Score 455.2; DB 9; Length 148828;
Best Local Similarity 96.8%; Pred.No. 4.4e-97;
Matches 486; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

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Qy 61 ATGGATACATATTGGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
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Qy 121 CAACGAGGATAGACACCGGAAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 180
Db 65206 CAACGAGGATAGACACCGGAAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 65265

Qy 181 GCAAGAGATTAAAGTTTTCATTTTGAAGCCATTGTGAATTTATACAAATAAAAAATG 240
Db 65266 GCAAGAGATTAAAGTTTTCATTTTGAAGCCATTGTGAATTTATACAAATAAAAAATG 65325

Qy 241 CAAGTGGATATTGAACAGTCTCTCTCTGATTAATTTCTGATACAGTACAGTTCACGCCCC 300
Db 65326 CAAGTGGATATTGAACAGTCTCTCTCTGATTAATTTCTGATACAGTACAGTTCACGCCCC 65385

Qy 301 TCACGAGACACTGAAACATGT -GGTCACCGCGAGACAGTGTGGCAATATTATCCCTGTAA 359
Db 65386 TCACGAGACACTGAAACATGTGGGTACCGGGAGACAGTGTGGCAATATTATCCCTGTAA 65445

Qy 360 TGTAACCAAGCTTGGCCAGAGCAGTGAACATATGACACAACTTTTGTGACAGTGGCTC 419
Db 65446 TGTAACCAAGCTTGGCCAAAGCAGTGAACATATGACACAACTTTTGTGACAGTGGCTC 65505

Qy 420 CTAATAGGACAGTCCGAGCCAAATTCAGCCAGTCCTTTCTGTGTTTATCCCATCTCTC 479
Db 65506 CTAATAGGACAGTCCGAGCCAAATTCAGCCAGTCCTTTCTGTGTTTATCCCATCTCTC 65565

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Db 65566 CCAATATTTGGAAACTGATGTCT 65589
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RESULT 8

AC090947/c

LOCUS

DEFINITION Homo sapiens chromosome 3 clone RP11-30G23 map 3p, complete sequence.

ACCESSION AC090947 AC016333

VERSION AC090947.2 GI:24796728

KEYWORDS HTG; HTGS DRAFT.

SOURCE Homo sapiens (human)

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 166049)

AUTHORS

Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J.,
Ding, H., Dong, W., Fan, H., Feng, J., Gong, J., Guan, Q., Gu, X., Guo, D.,
Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C.,
Li, F., Li, G., Li, J., Li, L., Li, S., Li, T., Liu, Y., Liu, N., Liu, B.,
Liu, Y., Li, W., Li, W., Li, Y., Luo, J., Luo, X., Qi, Q., Qi, X., Song, L.,
Song, S., Sun, M., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J.,

Wang, J., Wang, J., Wang, L., Wang, L., Wang, R., Wang, X., Wang, X.,
Wang, Y., Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B.,
Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M.,
Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N.,
Yu, J. and Yang, H.

Chromosome 3p genomic sequence

Unpublished

2 (bases 1 to 166049)

Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J.,
Ding, H., Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D.,
Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C.,
Li, F., Li, G., Li, J., Li, S., Li, T., Liu, Y., Liu, N., Liu, B.,
Liu, Y., Li, W., Li, W., Li, Y., Luo, J., Niu, Y., Qi, Q., Qi, X., Song, L.,
Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J.,
Wang, J., Wang, L., Wang, L., Wang, L., Wang, R., Wang, X., Wang, X.,
Wang, Y., Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B.,
Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M.,
Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N.,
Yu, J. and Yang, H.

Direct Submission

Submitted (20-MAR-2001) Human Genomic Center, Institute of
Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing
100101, P.R.China

3 (bases 1 to 166049)

Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J.,
Ding, H., Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D.,
Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C.,
Li, F., Li, G., Li, J., Li, S., Li, T., Liu, Y., Liu, N., Liu, B.,
Liu, Y., Li, W., Li, W., Li, Y., Luo, J., Niu, Y., Qi, Q., Qi, X., Song, L.,
Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J.,
Wang, J., Wang, L., Wang, L., Wang, L., Wang, R., Wang, X., Wang, X.,
Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B., Zeng, Y.,
Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M., Zhang, X.,
Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N., Yu, J. and
Yang, H.

Direct Submission

Submitted (08-NOV-2002) Human Genomic Center, Institute of
Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing
100101, P.R.China

On Nov 8, 2002 this sequence version replaced gi:13384351.

-----Genome Center

Center:Beijing Center

Center code:Beijing

Website: <http://hgsc.igtp.ac.cn><http://www.genomics.org.cn>Contact: hgsc@igtp.ac.cn

----- Project Information

Center project name: 13 project

Center clone name: RP11-30G23

----- Summary Statistics

Sequencing vector: pUC18; 100% of reads

Chemistry: Dye-terminator; ET 55% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 499 bases at least Q40

Consensus quality: 605 bases at least Q30

Consensus quality: 674 bases at least Q20

Insert size: 692; sum-of-contigs

Quality coverage: 2.80x in Q20 bases; sum-of-contigs

FEATURES

source

Location/Qualifiers
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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="3p"
/map="RP11-30G23"

ORIGIN

Query Match 90.5%; Score 455.2; DB 9; Length 166049;
Best Local Similarity 96.8%; Pred. No. 4.4e-97;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

Oy 1 TAGAGTAACTACCTTAGGAATATATACATTTTCAGTAGCATGCTGATACCAACGTTTAAACT 60
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145676 CAACAGGAGATAGACACCGGAAAGAAACTTTGGCCCAATAAGCTTTCTGGTATTTCTATAA 145617
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145496 TCACGAGACACTGAACATGTGGGTACCGCGGACAGTGTGGCAATATTTATCCCTGTAA 145437
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145436 TGTACCAAGTCTTCCAGAGAGAGTGAACATTTATGACACAACTTTTGTGACAGCTGGCTC 145377
Db
Oy 420 CTATAGGACAGTGTCCAGCCCAATTTCAAGCCAGTCTTCTGCTGTTTATTTCCCATCTCTC 479
145376 CTATAGGACAGTGTCCAGCCCAATTTCAAGCCAGTCTTCTGCTGTTTATTTCCCATCTCTC 145317
Db
Oy 480 CCAATATTTGGAACATGATGCT 503
145316 CCAATATTTGGAACATGATGCT 145293
Db

RESULT 9

AC151849/c

LOCUS

DEFINITION

ordered pieces.

AC151849

AC151849.1

GI:53983836

HTG; HTGS PHASE2; HTGS DRAFT.

KEYWORDS

Callithrix jacchus (white-tufted-ear marmoset)

SOURCE

Callithrix jacchus

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Rutheria; Primates; Platyrrhini; Callitrichidae;

Callithrix.

REFERENCE

1 (bases 1 to 186028)

AUTHORS

Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

Direct Submission

Unpublished

JOURNAL

2 (bases 1 to 186028)

REFERENCE

Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

Direct Submission

Submitted (08-OCT-2004) Genome Sciences, Lawrence Berkeley National

Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA

COMMENT

Sequence Produced by Berkeley PGA

Web site: <http://pga.lbl.gov>

Center Code: PGABERK

Center Project Name: J108-42F14

Bac Clone Name: CH259-42F14

This sequence has been compared to sequences of other species
using VISTA (<http://www-gad.lbl.gov/VISTA>). The results can be
viewed at:
http://pga.lbl.gov/cgi-bin/search_cvcgd?type=n&value=PPARG


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Db 15463 AAGATATGTATATATTTGAATACAAAGAGTATTTCTTCAGATAATGTGATTA-----AGA 15410
QY 117 GATTCAACAGGAGTAGACACCGAAGAAAACCTTTGGCCCAATAAGCTT-TCTGTATTT 175
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QY 294 ACGCCCTTCACGAGACACTGCAACATGTGTGTACCGGAGAGACAGTGTGGCAATATTATCC 353
Db 15229 ACACCCCTCACAACACTGAATGTGTGGTCACTGGCGAGACAGTGTAGCAACGTTTTTCC 15170
QY 354 CTGTAATGTACCAAGTCTTTGCCA---GAGCAGTGAACATTTATGACACAACTTTTGTGCAC 410
Db 15169 TTGTAATGTACCAAGTCTTTGCCAAGACGACAGACATTTATGACACACCACTTTTGTGCAC 15110
QY 411 AGCTGGCTCTTAAT-AGCAGAGTGCACGCCCAATTCACGCCAGTCTTCTGTGTATTT 469
Db 15109 AGCTGGCTCTCAATCAGGACAGTGCACGCCCAATTCAGGCCCTGTATCTTCTGTGTATTT 15050
QY 470 CCCATCTCTCCCAATATTTTGAACACTGATGTCT 503
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RESULT 12
AC120668/c
LOCUS
DEFINITION
Rattus norvegicus clone CH230-24K10, *** SEQUENCING IN PROGRESS
***, 3 unordered pieces.
AC120668
VERSION
AC120668.4 GI:25188255
KEYWORDS
HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
SOURCE
Rattus norvegicus
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 250169)
Muzny,D.Marie., Metzker,M.Lee., Abramson,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Albrooks,S., Amin,A., Anguiano,B.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,K., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Derramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregiorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hoques,M.,
Hollins,B., Howells,S., Hulyk,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensuhewa,L., Loulseghe,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindaratne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,

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Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S., Parks,K.,
Nwakoelamh,O., Okwuonu,G., Olarunpunaagoo,A., Pal,S., Parks,C.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkoch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,B., Pu,L.-L.,
Puzos,M., Quiroz,J., Rachlin,B., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savary,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Sivartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Soza,J.,
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 250169)
Worley,K.C.
Direct Submission
Submitted (09-MAY-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 250169)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (23-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

```

COMMENT

```

On Nov 23, 2002 this sequence version replaced gi:23322270.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GXLO
Center clone name: CH230-24K10
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 225414 bases at least Q40
Consensus quality: 228258 bases at least Q30
Consensus quality: 229994 bases at least Q20
Estimated insert size: 226717; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation
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* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence

```

* as soon as it is available and the accession number will
* be preserved.

1 9799: contig of 9799 bp in length
8900 9899: gap of unknown length
9900 248957: contig of 239058 bp in length
248958 249057: gap of unknown length
249058 250169: contig of 1112 bp in length.

FEATURES

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/note="wgs contig"
complement(234439..235077)
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site:EcoRI
end_sequence:BH280540"
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/note="wgs_end_extension
clone_end:T7"
247156..248957
/note="wgs_end_extension
clone_end:T7"

ORIGIN

Query Match 43.7%; Score 219.6; DB 2; Length 250169;
Best Local Similarity 75.3%; Pred. No. 1.3e-41;
Matches 342; Conservative 0; Mismatches 99; Indels 13; Gaps 5;
Qy 57 AACTATGATACATATTGTAATCCAAATTTCTTCAGATATGTAATAGAGATTAGA 116
Db 154629 AAGAATATGTATATTTGAATACAGAGTATCTTCAGATAATGTGATTA-----AGA 154576
Qy 117 GATTCAACCGGGATAGACACCGAAGAAACTTTGCCAAATAAGCTT-TCTGGTATTT 175
Db 154575 ATTTCAACCAAGGATGATACCTTAAGAAATGTTGCCAAATGAGTTTATCTAGTGT 154516
Qy 176 CATAGCAAGAGATTTAAGTTTTCATTTAAGAA--GCCATTGTGAATTTATACAAATA 233
Db 154515 CATAACTTAAAGATTTAAAGTTTCTATTAAAAAGCCACTGGTGTGTTTACTGCAA 154456
Qy 234 AAAAATGCAAGTATATGACAGTCTCTCTCTGATATTTCTTAATACAGTACAGTTC 293
Db 154455 TTAATAAGCAATCAATATTAACAGTCTCTCTCTGGTAATTCAAATCTGTACAGTTC 154396
Qy 294 ACGCCCTCAGGACACTGAACATGTGGTCACCGCGAGACAGTGTGGCAATATTATCC 353
Db 154395 ACACCCCTCAAAACACTGAATGTGTGGTCACTTGGCGAGACAGTGTAGCAACGTTTCC 154336
Qy 354 CTGTAATGTACCAAGTCTTTGCCA--GAGCAGTGAACATTTATGACAACTTTTGTGAC 410
Db 154335 TTGTAATGTACCAAGTCTTTGCCAAGAAGACAGACAGCATTTATGACACACCATTTTGTGCAC 154276
Qy 411 AGCTGGCTCTTAAT-AGGACAGTGCAGCCAAATTCAGCCAGTCCCTTCTGTGTTTATT 469
Db 154275 AGCTGGCTCTCAATCAGACAGTGCAGCCAAATTCAGCCCTGATCCCTTCTGTGTTTATT 154216
Qy 470 CCCATCTCTCCCAATATTTGGAACTGATGTCT 503
Db 154215 CCCACCTCTCCCAATATTTGAAACTGGTGTCT 154182

RESULT 13

RESULT 14
AY243584

S79407
LOCUS S79407 682 bp DNA linear ROD 09-DEC-1995
DEFINITION mppar gamma 2=peroxisome proliferator-activated receptor gamma isoform 2 [promoter] [mice, Genomic, 682 nt].
S79407
ACCESSION S79407.1 GI:1110564
VERSION S79407.1
KEYWORDS
SOURCE Mus sp.
ORGANISM Mus sp.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 682)
AUTHORS Zhu,Y., Qi,C., Korenberg,J.R., Chen,X.N., Noya,D., Rao,M.S. and Reddy,J.K.
TITLE Structural organization of mouse peroxisome proliferator-activated receptor gamma (mppar gamma) gene: alternative promoter use and different splicing yield two mppar gamma isoforms
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 92 (17), 7921-7925 (1995)
MEDLINE 95372391
PUBMED 7644514
REMARK GenBank staff at the National Library of Medicine created this entry [NCBI gbbseq 170483] from the original journal article.
FEATURES
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616..>682
680..682
/genes="mppar<gamma>2"
/notes="peroxisome proliferator-activated receptor gamma isoform 2"

ORIGIN

Query Match 43.0%; Score 216.4; DB 10; Length 682;
Best Local Similarity 77.6%; Pred. No. 1.3e-40;
Matches 354; Conservative 0; Mismatches 86; Indels 16; Gaps 7;
Qy 57 AACTATGATACATATTGTAATCCAAATTTCTTCAGATATGTAATAGAGATTAGA 116
Db 108 AAGAATGTGTATATGTTTGTAGTACAGAAATATTTCTTCAGAT-GTGTGATTAG-----GA 160
Qy 117 GATTCAACCGGGATAGACACCGAAGAAACTTTGCCAAATAAGCTT-TCTGGTATTT 175
Db 161 GTTTCACCAAGATAATACTTAAGAAAACTTTGCCAAATACGTTTATCTGTGTTT 220
Qy 176 CATAAGCAAGAGATTTAAGTTTTCATTTAAGAACGCA---TTGTGAATTATACAAAT 232
Db 221 CATAACTTAGAGATTAAGGTTTCTATTTTAAAAAGCCACTGGTGTGTTTACTGCAAT 280
Qy 233 AAAAATGCAAGTGGATTTGAACAGTCTCTCTCTGATATTTCTAAATACAGTACAGTT 292
Db 281 TTTAAAAAGCAATCAATTTTGAACATCTCTCTCTGGTAATTCCAACTACTGTACAGTT 340
Qy 293 CAGCCCCCTCAGGACACATGAACATGT-GGTCAACCGCGAGACAGTGTGGCAATATTAT 351
Db 341 CAGCCCCCTCAGACAGACAGTGAATGTGTGGTCACTGGCGAGACAAATGTAGCAACGTTT 400
Qy 352 CCCTGTAATGTACCAAGTCTTTGCCA---GAGCAGTGAACATTTATGACACACTTTTGTGTC 408
Db 401 CCTTGTAAATGTACCAAGTCTTGGCAAGACAGACAGCATTTATGACACACATTTTGTGTC 460
Qy 409 ACAGCTGGCTCTTAAT-AGGACAGTGCAGCCAAATTCAGCCAGTCTCTTCTGTGTTTA 467
Db 461 ACAAATGGCTCTCAATCAGACAGTGCAGCCAAATTCAGCCCTGATTTCTTCTGTGTTTA 520
Qy 468 TTCCCATCTCTCCCAATATTTGGAAACTGATGTCT 503
Db 521 TTCCCATCTCTCCCAATATTTGAAAACTGGTGTCT 556

LOCUS AY243584 1159 bp DNA linear ROD 01-JUL-2003
DEFINITION Mus musculus strain 129S1/SvImJ peroxisome proliferator-activated
receptor gamma transcript 2 (Pparg) gene, promoter region.
ACCESSION AY243584
VERSION AY243584.1 GI:32395962
KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 1159)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE Lith13 Encompasses Lrpap1, Which is a Likely Genetic Determinant of
Murine Cholesterol Gallstone Formation
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1159)
AUTHORS Lyons, M.A., Wittenburg, H., Li, R., Walsh, K.A., Leonard, M.R.,
Churchill, G.A., Carey, M.C. and Paigen, B.
TITLE Lith13 Encompasses Lrpap1, Which is a Likely Genetic Determinant of
Murine Cholesterol Gallstone Formation
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1159)
AUTHORS Lyons, M.A., Wittenburg, H., Walsh, K.A., Carey, M.C. and Paigen, B.
TITLE Direct Submission
JOURNAL Submitted (25-FEB-2003) The Jackson Laboratory, 600 Main Street,
Bar Harbor, ME 04609, USA
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Location/Qualifiers
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/note="peroxisome proliferator-activated receptor gamma
transcript 2"
1. .1159
/gene="Pparg"
gene
promoter
ORIGIN
Query Match 43.0%; Score 216.4; DB 10; Length 1159;
Best Local Similarity 77.6%; Pred. No. 1.2e-40;
Matches 354; Conservative 0; Mismatches 86; Indels 16; Gaps 7;
Qy 57 AACTATGATACATATTGTAATTCCTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGA 116
Db 633 AAGAATGTGTATATGTTTGAGTACAGAATATTTCTTCAGAT-GTGTGATTAG-----GA 685
Qy 117 GATTCAACCGGGATAGACACCGAAGAAACTTTGCCAAATAAGCTT-TCTGGTATTT 175
Db 686 GTTTCACCAAGATAATATCTTAAGRAAAACTTTGGCCAAATACGTTTATCTGGTGT 745
Qy 176 CATAGCAAGAGATTTTAAGTTTTCATTTAAGAAGCCA---TTGTGAATTATACAAAT 232
Db 746 CATAACTTAGAGATTAAGTTTCTATTTTAAAGCCACTGGTGTGTTTACTGCAAT 805
Qy 233 AAAAAATGCAAGTGGATTTGAACAGTCTCTCTCTGATTAATTTCTTAATACAGTACAGTT 292
Db 806 TTTAAAAAGCAATCAATATTGAACAACTCTGCTCTGTTAATTCCACTACTGACAGTT 865
Qy 293 CACGCCCTCACGAGACACTGAACATGT-GGTCAACCGCGAGACAGTGTGGCAATATTAT 351
Db 866 CACGCCCTCACAGACAGTGAATGTGTGGGTCACTGGCGAGACATGTAGCAAGTTT 925
Qy 352 CCCTGTAATGTACCAAGCTTTGCCA---GAGCAGTGAACATTTATGACACAACTTTTGTGTC 408
Db 926 CCTTGTAAATGTACCAAGCTTTGCCAAGCAGTGAATGTGTGGGTCACTGGCGAGACATGTAGCAAGTTT 985
Qy 409 ACAGCTGGCTCTTAAT-AGGACAGTGGCCCAATTCAGCCAGTCTCTTCTCTGTGTTA 467
Db 986 ACAACTGGCTCTCAGTCAGGACAGTGGCCCAATTCAGCCAGTCTCTTCTCTGTGTTA 1045
Qy 468 TTCCCATCTCTCCCAAAATTTTGAAGAACTGATGCT 503
Db 1046 TTCCCATCTCTCCCAAAATTTTGAAGAACTGATGCT 1081

RESULT 15
AY236531 1160 bp DNA linear ROD 01-JUL-2003
LOCUS AY236531
DEFINITION Mus musculus strain DBA/2J peroxisome proliferator-activated
receptor gamma transcript 2 (Pparg) gene, promoter region.
ACCESSION AY236531
VERSION AY236531.1 GI:32395932
KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 1160)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE Lith6 Encompasses Ppargamma and SLC21A1 Which Are Likely Genetic
Determinants of Murine Cholesterol Gallstone Formation
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1160)
AUTHORS Lyons, M.A., Wittenburg, H., Walsh, K.A., Leonard, M.R.,
Korstanje, R., Churchill, G.A., Carey, M.C. and Paigen, B.
TITLE Lith6 Encompasses Ppargamma and SLC21A1 Which Are Likely Genetic
Determinants of Murine Cholesterol Gallstone Formation
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1160)
AUTHORS Lyons, M.A., Wittenburg, H., Walsh, K.A., Carey, M.C. and Paigen, B.
TITLE Direct Submission
JOURNAL Submitted (13-FEB-2003) The Jackson Laboratory, 600 Main Street,
Bar Harbor, ME 04609, USA
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1. .>1160
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transcript 2"
1. .1160
/gene="Pparg"
gene
promoter
ORIGIN
Query Match 43.0%; Score 216.4; DB 10; Length 1160;
Best Local Similarity 77.6%; Pred. No. 1.2e-40;
Matches 354; Conservative 0; Mismatches 86; Indels 16; Gaps 7;
Qy 57 AACTATGATACATATTGTAATTCCTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGA 116
Db 634 AAGAATGTGTATATGTTTGAGTACAGAATATTTCTTCAGAT-GTGTGATTAG-----GA 686
Qy 117 GATTCAACCGGGATAGACACCGAAGAAACTTTGCCAAATAAGCTT-TCTGGTATTT 175
Db 687 GTTTCACCAAGATAATATCTTAAGRAAAACTTTGGCCAAATACGTTTATCTGGTGT 746
Qy 176 CATAGCAAGAGATTTTAAGTTTTCATTTAAGAAGCCA---TTGTGAATTATACAAAT 232
Db 747 CATAACTTAGAGATTAAGTTTCTATTTTAAAGCCACTGGTGTGTTTACTGCAAT 806
Qy 233 AAAAAATGCAAGTGGATTTGAACAGTCTCTCTCTGATTAATTTCTTAATACAGTACAGTT 292
Db 807 TTTAAAAAGCAATCAATATTGAACAACTCTGCTCTGTTAATTCCACTACTGACAGTT 866
Qy 293 CACGCCCTCACGAGACACTGAACATGT-GGTCAACCGCGAGACAGTGTGGCAATATTAT 351
Db 867 CACGCCCTCACAGACAGTGAATGTGTGGGTCACTGGCGAGACATGTAGCAAGTTT 926
Qy 352 CCCTGTAATGTACCAAGCTTTGCCA---GAGCAGTGAACATTTATGACACAACTTTTGTGTC 408
Db 927 CCTTGTAAATGTACCAAGCTTTGCCAAGCAGTGAATGTGTGGGTCACTGGCGAGACATGTAGCAAGTTT 986
Qy 409 ACAGCTGGCTCTTAAT-AGGACAGTGGCCCAATTCAGCCAGTCTCTTCTCTGTGTTA 467
Db 987 ACAACTGGCTCTCAGTCAGGACAGTGGCCCAATTCAGCCAGTCTCTTCTCTGTGTTA 1046
Qy 468 TTCCCATCTCTCCCAAAATTTTGAAGAACTGATGCT 503

Db 1047 TTCCACCTCTCCCAATATTTGAAACTGGTGCT 1082

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Job time : 2644.94 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:46:48 ; Search time 407.412 Seconds
(without alignments)
7308.644 Million cell updates/sec

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Sequence: 1 tagagtaagtaccttagaa.....atatattgaaactgatgtct 503

Scoring table: IDENTITY NUC
Gapop 10'0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 1: Geneseqn1980s:*
- 2: Geneseqn1990s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004as:*
- 13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	503	100.0	2045	2	AX19033 Human PPA
2	455.2	90.5	1100	5	Aaf23651 Human PPA
C 3	455.2	90.5	85873	10	ADH10008 Human chr
4	455.2	90.5	158417	13	AdS36461 Human aut
C 5	455.2	90.5	166043	12	AdL08127 Human gen
6	216.4	43.0	1184	10	ACC79611 Mouse per
7	52	10.3	177	2	AX19064 Human PPA
8	45.2	9.0	200000	12	Ado47191 DNA seque
C 9	44.2	8.8	3162	6	Abq72637 Human MDD
C 10	43.6	8.7	6215	6	AbL33191 Human imm
C 11	41.8	8.3	99916	6	Adi03931 Human enz
C 12	41.8	8.3	116297	12	Adq97587 Human can
C 13	41.2	8.2	32392	6	AbL56203 AMEPV gen
C 14	40.8	8.1	2000	12	AdJ40655 Plant cdn
C 15	40.8	8.1	109661	12	Adq97818 Human can
C 16	40.2	8.0	376	4	Aah93354 Human chr
C 17	40.2	8.0	2000	8	Ada71938 Rice gene
C 18	40	8.0	8056	8	AbZ10100 Haematopo
19	39.4	7.8	6191	6	AbL33216 Human imm
20	39.4	7.8	6191	6	AbK31306 Signal tr

21	39.4	7.8	6191	6	ABL70281	ABL70281 Chemical
22	39.4	7.8	6191	6	ABN80160	ABN80160 Human che
C 23	39.2	7.8	6308	6	ABL33470	ABL33470 Human imm
C 24	39	7.8	6565	4	AS46466	AS46466 Tumour su
C 25	39	7.8	6565	6	ABK31327	ABK31327 Signal tr
C 26	39	7.8	37515	6	ABQ66998	ABQ66998 Human ang
27	38.8	7.7	1696	12	ADI43067	ADI43067 Plant tra
28	38.8	7.7	1696	12	ADO03207	ADO03207 Corn orth
C 29	38.8	7.7	8056	8	ABZ10246	ABZ10246 Haematopo
C 30	38.8	7.7	10945	6	ABK28354	ABK28354 DNA trans
C 31	38.8	7.7	19634	8	ABZ10162	ABZ10162 Haematopo
C 32	38.8	7.7	19634	8	ABZ10016	ABZ10016 Haematopo
C 33	38.8	7.7	19634	13	ADS89398	ADS89398 Oligonuc
C 34	38.8	7.7	19634	13	ADS89672	ADS89672 Oligonuc
C 35	38.4	7.6	2258	4	ABL21792	ABL21792 Drosophil
C 36	38.2	7.6	4152	6	ABL55640	ABL55640 AMEPV ABC
C 37	38.2	7.6	50000	6	ABL56201	ABL56201 AMEPV gen
C 38	38.2	7.6	96589	9	ADA02708	ADA02708 Human ZFH
C 39	38.2	7.6	96589	10	ADB72446	ADB72446 Human ZFH
C 40	38.2	7.6	96589	10	ADE95956	ADE95956 Human ZFH
41	38	7.6	2035	13	ADT05376	ADT05376 Haemophil
C 42	38	7.6	349980	13	ADT05649	ADT05649 Haemophil
43	37.8	7.5	2142	2	AX84309	AX84309 Stealth v
44	37.8	7.5	2144	3	AAZ36928	AAZ36928 Nucleotid
C 45	37.8	7.5	5274	11	ACN92690	ACN92690 Breast ca

ALIGNMENTS

RESULT 1

AX19033
ID AX19033 standard; DNA; 2045 BP.

AC AX19033;

DT 13-MAY-1999 (first entry)

XX Human PPAR-gamma-2 proximal promoter, exon B and intron B.

Human; peroxisome proliferator activated receptor gamma; PPAR-gamma; regulatory sequence; promoter; obesity; anorexia; lipoma; cachexia; lipodystrophy; liposarcoma; human immunodeficiency virus; HIV; insulin resistance; non-insulin-dependent diabetes mellitus; polycystic ovary syndrome; gastrointestinal tract; Crohn's disease; inflammatory bowel disease; ulcerative colitis; bowel cancer; ss.

OS Homo sapiens.

XX WO9905161-A1.

XX 04-FEB-1999.

XX 24-JUL-1998; 98WO-US015411.

XX 25-JUL-1997; 97US-0053692P.

XX (LIGA-) LIGAND PHARM INC.

XX (INSP) INST PASTEUR.

XX Briggs MR, Saladin RS, Auwerx J, Fajas L;

XX WPI, 1999-142844/12.

XX Newly isolated nucleic acid comprising a control region of a human peroxisome proliferator activated receptor (PPAR) gamma gene - useful for identifying modulators that are useful in treating diseases associated with abnormal levels of human PPAR-gamma gene expression.

XX Claim 10; Page 80; 102pp; English.

XX The present invention describes an isolated, purified or enriched nucleic acid comprising a control region of a human peroxisome proliferator

activated receptor gamma (PPAR-gamma) gene. The nucleic acids are useful for screening for agents capable of modulating the expression of a human PPAR-gamma gene. These agents (modulators) form pharmaceutical compositions that are useful for treating diseases associated with high/low levels of human PPAR-gamma gene expression. The diseases include obesity, anorexia, cachexia, lipodystrophy, lipomas, liposarcomas, abnormalities associated with anti-human immunodeficiency virus (HIV) treatment, insulin resistance, non-insulin-dependent diabetes mellitus (NIDDM), polycystic ovary syndrome, diseases of the gastrointestinal (GI) tract, inflammatory bowel disease, Crohn's disease, ulcerative colitis and bowel cancer. The nucleic acids are useful for studying the role of the PPAR-gamma gene in various diseases and disorders. The structure of PPAR-gamma enables genetic studies of PPAR-gamma mutations in humans, and evaluation of its role in disorders like insulin resistance, NIDDM, and diseases associated with altered adipose tissue function, like obesity and lipodystrophic syndromes. The nucleic acids are also useful for gene therapy and the production of transgenic animals, which are useful in screening assays. The control regions of the nucleic acids enable screening for modulators of the human PPAR-gamma gene, which are useful in designing drugs for treating disorders or diseases associated with the level of PPAR-gamma gene expression. The present sequence represents human PPAR-gamma-2 proximal promoter, exon B and intron B

XX SQ Sequence 2045 BP; 605 A; 422 C; 351 G; 667 T; 0 U; 0 Other;

Query Match 100.0%; Score 503; DB 2; Length 2045;
Best Local Similarity 100.0%; Pred. No. 2.5e-128;
Matches 503; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAAACT 60
Db 818 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAAACT 877
Qy 61 ATGATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
Db 878 ATGATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 937
Qy 121 CAACACGGGATAGACACCGGAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 180
Db 938 CAACACGGGATAGACACCGGAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 997
Qy 181 GCAAGAGATTAAAGTTTCCATTTTAAGAGCCATTTGTAATATATACCAATAAAAAATG 240
Db 998 GCAAGAGATTAAAGTTTCCATTTTAAGAGCCATTTGTAATATATACCAATAAAAAATG 1057
Qy 241 CAAGTGGATATTGACAGTCTCTTCTGTGATAATTTCTAAATACAGTACAGTTTCACGCC 300
Db 1058 CAAGTGGATATTGACAGTCTCTTCTGTGATAATTTCTAAATACAGTACAGTTTCACGCC 1117
Qy 301 TCACGAGACACTGAAACATGTGGTCAACCGGCGAGACAGTGGCCAAATATTATCCCTGTAAT 360
Db 1118 TCACGAGACACTGAAACATGTGGTCAACCGGCGAGACAGTGGCCAAATATTATCCCTGTAAT 1177
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Db 1178 GTACAAGTCTTGGCAGAGCAGTGAACATTAATGACAACTTTTGTGACAGTGGCTCC 1237
Qy 421 TAATAGGACAGTGGCCCAATTCAGCCAGTCTCTTCTGTGTTTATTTCCCATCTCTCC 480
Db 1238 TAATAGGACAGTGGCCCAATTCAGCCAGTCTCTTCTGTGTTTATTTCCCATCTCTCC 480
Qy 481 CAAATATTGGAAACTGATGTCT 503
Db 1298 CAAATATTGGAAACTGATGTCT 1320

RESULT 2

AAF23651

ID AAF23651 standard; cDNA; 1100 BP.

XX

AC AAF23651;

XX

DT 27-MAR-2001 (first entry)

XX Human PPARgamma N-terminal sequence coding sequence.
DS
XX Cytostatic; antiinflammatory; antisense oligonucleotide; PPARgamma;
KW peroxisome proliferator-activated receptor gamma; transcription factor;
KW nuclear hormone receptor; human; infection; inflammation; tumour; ss.
XX Homo sapiens.
XX OS
XX US6159734-A.
XX 12-DEC-2000.
XX 18-JAN-2000; 2000US-00484345.
XX 18-JAN-2000; 2000US-00484345.
XX (ISIS-) ISIS PHARM INC.
XX McKay R, Baker BF, Borchers AH;
XX WPI; 2001-070112/08.
XX P-PSDB; AAB59840.
XX Novel antisense compounds capable of modulating expression of peroxisome
PT proliferator-activated receptor gamma useful for diagnosis, prophylaxis
PT and treatment of diseases associated with expression of the receptor.
XX Example 15; Col 49-52; 40pp; English.
XX Peroxisome proliferator-activated receptors (PPARs) are members of the
CC nuclear hormone receptor subfamily of transcription factors. The present
CC invention relates to antisense oligonucleotides (see AAF23652-P23731),
CC targeted to a nucleic acid molecule encoding human PPARgamma, which
CC specifically hybridises with and inhibits the expression of human
CC PPARgamma. The present sequence is the coding sequence for the N-terminal
CC sequence of human PPARgamma. The PPARgamma antisense oligonucleotides of
CC the present invention can be used in the diagnosis and treatment of
CC diseases associated with the expression of PPARgamma, e.g. to prevent or
CC delay infection, inflammation or tumour formation
XX SQ Sequence 1100 BP; 334 A; 215 C; 195 G; 356 T; 0 U; 0 Other;

Query Match 90.5%; Score 455.2; DB 5; Length 1100;
Best Local Similarity 96.8%; Pred. No. 3.1e-115;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

Qy 1 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAAACT 60
Db 239 TAGAGTAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAAACT 298
Qy 61 ATGATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
Db 299 ATGATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATT 351
Qy 121 CAACACGGGATAGACACCGGAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 180
Db 352 CAACACGGGATAGACACCGGAAGAAACTTTGCCCAATAAGCTTTCTGGTATTTCATAA 411
Qy 181 GCAAGAGATTAAAGTTTTCATTTTAAGAGCCATTTGTAATATATACCAATAAAAAATG 240
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Db 472 CAAAGTGGATATTGAACAGTCTCTTCTGTGATAATTTCTAAATACAGTACAGTTTCACGCC 531
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Db 532 TCACGAGACACTGAACATGTGGGTACCGCGGAGACAGTGTGGCAATATTATTCCTGTAA 591
Qy 360 TGTACCAAGTCTTGGCCAGAGCAGTGAACATTTATGACACCAACTTTTCTCAGCTGCTC 419

Db 592 TGTACCAAGTCTTGCCAAAGCAGTGAACATTTATGACACAACTTTTGTGCAGCTGGCTC 651
Qy 420 CTAATAGACAGTGCAGCCAAATTCAGCCAGTCTTCTGTGTATTCCTCCATCTCTC 479
Db 652 CTAATAGACAGTGCAGCCAAATTCAGCCAGTCTTCTGTGTATTCCTCCATCTCTC 711
Qy 480 CCAATATTTGGAACACTGATGCT 503
Db 712 CCAATATTTGGAACACTGATGCT 735

RESULT 3

ADH10008/c

ID ADH10008 standard; DNA; 85873 BP.

XX AC

XX ADH10008;

XX DT

XX 11-MAR-2004 (first entry)

XX XX

XX Human chromosome 3p25 DNA fragment.

XX KW

XX ds; human; chromosome 2; hyperplasia; tumour; thyroid; cancer;

XX KW break point; chromosomal band 2p21-22; DRIP; cytostatic; thyromimetic;

XX KW chromosome 7p15.

XX XX

XX Synthetic.

XX OS

XX Homo sapiens.

XX XX

XX WO2003093310-A1.

XX PN

XX 13-NOV-2003.

XX PD

XX 02-MAY-2003; 2003WO-EP004642.

XX XX

XX 01-MAY-2002; 2002DE-01019413.

XX PR

XX 14-SEP-2002; 2002DE-01042705.

XX XX

XX (UYBR-) UNIV BREMEN.

XX PA

XX Bullerdiel J;

XX PI

XX WPI; 2003-854480/79.

XX DR

XX New nucleic acid from human chromosome 2, useful for treatment and

XX PT diagnosis of thyroid disease, especially cancer, also related

XX PT polypeptides and modulators.

XX XX

XX Claim 1; SEQ ID NO 3; 461pp; German.

XX PS

XX This invention describes a novel nucleic acid from human chromosome 2

XX CC that shows altered expression in hyperplasia and/or tumours, especially

XX CC of the thyroid. The invention describes a method for preparing nucleic

XX CC acid that can be detected in thyroid cancers where these contain an

XX CC aberration of the break point in chromosomal band 2p21-22. The invention

XX CC also describes constructs comprising a fragment of the human DRIP gene

XX CC and its splice variants, one of which contains all of exons 1-38 while

XX CC the other lacks exons 27 and 28. Some tumours contain a fusion protein of

XX CC DRIP, including a small segment from chromosome 3. The products of the

XX CC invention have cytostatic and thyromimetic activity and are used for the

XX CC inhibition of nucleic acid expression by antisense, ribozyme or RNA

XX CC interference (RNAi) methods. This sequence represents a fragment of human

XX CC chromosome 3p15 near the region encoding the DRIP protein described in

XX CC the disclosure of the invention.

XX SQ

SQ Sequence 85873 BP; 27329 A; 16422 C; 15988 G; 26134 T; 0 U; 0 Other;

XX Query Match

XX Best Local Similarity 90.5%; Score 455.2; DB 10; Length 85873;

XX Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

XX 1 TAGAGTAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAACT 60

XX Qy

XX 54472 TAGAGTAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATACCAACGTTTAACT 54413

XX Db

Qy 61 ATGGATACATATTTGAATTTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
Db 54412 ATGGATACATATTTGAATTTCCAAATTTTCTTCAAATTAATGT-----GATTAGAGATT 54360
Qy 121 CAACAGGAGATAGACACGAAAGAAACTTTGGCCAAATAAGCTTTCTGGTATTTTCATAA 180
Db 54359 CAACAGGAGATAGACACGAAAGAAACTTTGGCCAAATAAGCTTTCTGGTATTTTCATAA 54300
Qy 181 GCAAGAGATTTAAGTTTTCATTTTAAAGAGCCATTTGTGAATTTATACACAATAAAAAATG 240
Db 54299 GCAAGAGATTTAAGTTTTCATTTTAAAGAGCAATTTGTGAATTTTACAACAATAAAAAATG 54240
Qy 241 CAAGTGGATATTTGAACAGTCTCTCTCTCTGATAATTTCTAAATACAGTACAGTTTCACGCCCC 300
Db 54239 CAAGTGGATATTTGAACAGTCTCTCTCTCTGATAATTTCTAAATACAGTACAGTTTCACGCCCC 54180
Qy 301 TCACGAGACACTGAACATGT--GGTCAACGGGAGACAGTGTGGCAATATTTATCCCTGTAA 359
Db 54179 TCACAAGACACTGAACATGTGGGTCAACGGGAGACAGTGTGGCAATATTTATCCCTGTAA 54120
Qy 360 TGTACCAAGTCTTCCAGAGCAGTGAACATTTATGACACAACCTTTTGTGCAGAGCTGGCTC 419
Db 54119 TGTACCAAGTCTTCCAGAGCAGTGAACATTTATGACACAACCTTTTGTGCAGAGCTGGCTC 54060
Qy 420 CTAATAGGACAGTGCAGCCCAATTTCAAGCCAGTCTCTTCTGTGTATTTCCCATCTCTC 479
Db 54059 CTAATAGGACAGTGCAGCCCAATTTCAAGCCAGTCTCTTCTGTGTATTTCCCATCTCTC 54000
Qy 480 CCAATATTTTGGAAACTGATGCT 503
Db 53999 CCAATATTTTGGAAACTGATGCT 53976

RESULT 4

ADS36461

ID ADS36461 standard; DNA; 158417 BP.

XX AC

XX ADS36461;

XX XX

XX 16-DEC-2004 (first entry)

XX DT

XX Human autoimmune disease-related genomic DNA sequence - SEQ ID 1675.

XX DE

XX single nucleotide polymorphism detection; SNP detection;

XX KW rheumatoid arthritis; type 1 diabetes; multiple sclerosis;

XX KW systemic lupus erythematosus; inflammatory bowel disease; psoriasis;

XX KW thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;

XX KW glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;

XX KW primary systemic vasculitis; ds.

XX OS

XX Homo sapiens.

XX XX

XX WO2004083403-A2.

XX PN

XX 30-SEP-2004.

XX PD

XX 18-MAR-2004; 2004WO-US008461.

XX PF

XX 18-MAR-2003; 2003US-0455444P.

XX XX

XX 25-APR-2003; 2003US-0465241P.

XX PR

XX (APPL-) APPLERA CORP.

XX XX

XX Cargill M, Begovich AB, Alexander HC;

XX PI

XX WPI; 2004-728480/71.

XX XX

XX New isolated nucleic acid molecule comprises at least 8 contiguous

XX PT nucleotides where one of the nucleotides is a single nucleotide

XX PT polymorphism (SNP), useful for diagnosing or treating autoimmune

XX PT diseases, e.g. rheumatoid arthritis.

XX XX

```
PS Claim 16; SEQ ID NO 1675; 123pp; English.
XX
CC The invention comprises amino acid and coding sequences containing
CC genetic polymorphisms associated with an altered risk of developing an
CC autoimmune disease (e.g. rheumatoid arthritis). The invention further
CC comprises a method of identifying an individual that has an altered risk
CC of developing an autoimmune disease, comprising detecting a single
CC nucleotide polymorphism (SNP) in a nucleic acid of the invention. The DNA
CC and protein sequences of the invention are useful for diagnosing and
CC treating autoimmune diseases, such as: rheumatoid arthritis, type 1
CC diabetes, multiple sclerosis, systemic lupus erythematosus, inflammatory
CC bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious
CC anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease,
CC myocarditis, Sjogren's disease, or primary systemic vasculitis. The
CC present nucleic acid represents a human autoimmune disease-related
CC genomic DNA sequence of the invention. NOTE: The present sequence is not
CC shown in the specification, but has been retrieved from the WIPO website.
XX
SQ Sequence 158417 BP; 47887 A; 30186 C; 31475 G; 48577 T; 0 U; 292 Other;

Query Match          90.5%; Score 455.2; DB 13; Length 158417;
Best Local Similarity 96.8%; Pred. No. 1.6e-114;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

Qy 1 TAGAGTAAGTACCTTAGGAATATAACATTTTCAGTAGCATGCTGTATACCAACGTTTAAACT 60
Db TAGAGTAAGTACCTTAGGAATATAACATTTTCAGTAGCATGCTGTATACCAACGTTTAAACT 69123

Qy 61 ATGATACATATTTGAATTCCAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
Db ATGATACATATTTGAATTCCAATTTTCTTCAGATAATGTGATTAGAGATT 69176

Qy 121 CAACCGGATAGACACCGAAGAAACTTTGCCAATAGCTTCTGCTATTTCATAA 180
Db CAACCGGATAGACACCGAAGAAACTTTGCCAATAGCTTCTGCTATTTCATAA 69236

Qy 181 GCAAGAGATTAAAGTTTCCATTTAAGAGCCATTGTGAATTTATACAAATAAAAAATG 240
Db GCAAGAGATTAAAGTTTCCATTTAAGAGCCATTGTGAATTTATACAAATAAAAAATG 69296

Qy 241 CAAGTGGATATGAACAGTCTCTCTCTGATAATTTCTAAATPACAGTTCACGCCCC 300
Db CAAGTGGATATGAACAGTCTCTCTCTGATAATTTCTAAATPACAGTTCACGCCCC 69356

Qy 301 TCACGAGACACTGAACATGT-GGTTCACCGGCGAGCAGTGTGGCAATATTATCCCTGTAA 359
Db TCACGAGACACTGAACATGTGGGTTCACCGGCGAGCAGTGTGGCAATATTATCCCTGTAA 69416

Qy 360 TGTACCAAGTCTTGCCAGAGCAGTGAACATTTATGACACAACTTTTGTGTCAGCTGGCTC 419
Db TGTACCAAGTCTTGCCAGAGCAGTGAACATTTATGACACAACTTTTGTGTCAGCTGGCTC 69476

Qy 420 CTAATAGACAGTGCAGCCAAATTCAGCCCAAGTCTTTCTGTGTTTATTCCTATCTCTC 479
Db CTAATAGACAGTGCAGCCAAATTCAGCCCAAGTCTTTCTGTGTTTATTCCTATCTCTC 69536

Qy 480 CCAATATTTTGGAACTGATGCT 503
Db CCAATATTTTGGAACTGATGCT 69560

RESULT 5
ADL08127/c
ID ADL08127 standard; DNA; 166043 BP.
AC ADL08127;
XX
XX 20-MAY-2004 (first entry)
XX
XX Human gene associated with low HDL-C PPARG.
XX
XX Human; ds; SNP; single nucleotide polymorphism;
XX high density lipoprotein-C; HDL-C; vascular disease; metabolic disease;
KW
```

```
KW coronary artery disease; gene.
XX Homo sapiens.
XX
XX Key Location/Qualifiers
FT variation /*tag= a
FT /standard_name= "Single nucleotide polymorphism"
XX
XX US2004043389-A1.
XX
XX 04-MAR-2004.
XX
XX 04-SEP-2002; 2002US-00235192.
XX
XX 04-SEP-2002; 2002US-00235192.
XX
XX (VITI-) VITIVITY INC.
XX
XX McCarthy J;
XX
XX WPI; 2004-214170/20.
XX
XX
XX Determining whether a subject has, or is at risk of developing, an
XX abnormally low high density lipoprotein-C (HDL-C) level comprises
XX detecting an allelic variant of a polymorphic region from any of a set of
XX 27 genes.
XX
XX Disclosure; SEQ ID NO 46; 37pp; English.
XX
XX The invention relates to determining whether a subject has, or is at risk
XX of developing, an abnormally low high density lipoprotein-C (HDL-C) level
XX comprises determining whether the subject has an allelic variant of a
XX polymorphic region from any of 27 genes (alleles listed in Table 5 of the
XX specification). Also included are determining whether a male subject has,
XX or is at risk of developing, an abnormally low HDL-C level, comprising
XX determining whether the male subject has an allelic variant of a
XX polymorphic region listed in Table 5 which is associated with abnormally
XX low HDL-C levels in males, and determining whether a female subject has,
XX or is at risk of developing, an abnormally low HDL-C level, comprising
XX determining whether the female subject has an allelic variant of a
XX polymorphic region listed in Table 5 which is associated with abnormally
XX low HDL-C levels in females. The allelic variant in determining whether a
XX subject has, or is at risk of developing, an abnormally low HDL-C level
XX is APOA 1 CC, CD14 1 CT, COL5A2 1 GG, EDNRB 1 AG or AA, FABP3 1 CT, GBE1
XX 1 AG or GG, LIPI 5 AA, MTHFR 1 CC, VWF 2 GG, or their complements. The
XX allelic variant in determining whether a male subject has, or is at risk
XX of developing, an abnormally low HDL-C level, LRPI 3 CC or CT, PAI2 4 GG,
XX 1 GG, CD14 1 CT or CC, and FABP3 1 CT, in combination, or their
XX complements. The methods are useful for diagnosing (a predisposition to)
XX abnormally low levels of low high density lipoprotein-C (HDL-C) in a
XX subject. The methods are useful in diagnosing (a predisposition to) or
XX prognosticating diseases and disorders associated with abnormal lipid
XX levels such as vascular and metabolic diseases, e.g., coronary artery
XX disease. The present sequence is a human gene containing a SNP (single
XX nucleotide polymorphism associated with low high density lipoprotein-C
XX (HDL-C) levels.
XX
XX Sequence 166043 BP; 48570 A; 35113 C; 34446 G; 47914 T; 0 U; 0 Other;

Query Match          90.5%; Score 455.2; DB 12; Length 166043;
Best Local Similarity 96.8%; Pred. No. 1.6e-114;
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;

Qy 1 TAGAGTAAGTACCTTAGGAATATAACATTTTCAGTAGCATGCTGTATACCAACGTTTAAACT 60
Db TAGAGTAAGTACCTTAGGAATATAACATTTTCAGTAGCATGCTGTATACCAACGTTTAAACT 145789

Qy 61 ATGATACATATTTGAATTCCAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATT 120
Db ATGATACATATTTGAATTCCAATTTTCTTCAGATAATGTGATTAGAGATT 145677
```


XX PD 04-FEB-1999.
XX PF 24-JUL-1998; 98WO-US015411.
XX PR 25-JUL-1997; 97US-0053692P.
XX PA (LIGA-) LIGAND PHARM INC.
XX PA (INSP) INST PASTEUR.
XX FI Briggs MR, Saladin RS, Auwerx J, Pajas L;
XX DR WPI; 1999-142844/12.
XX XX
XX PT Newly isolated nucleic acid comprising a control region of a human
XX PT peroxisome proliferator activated receptor (PPAR) gamma gene - useful for
XX PT identifying modulators that are useful in treating diseases associated
XX PT with abnormal levels of human PPAR-gamma gene expression.
XX XX
XX PS Disclosure; Page 87; 102pp; English.
XX XX
XX CC The present invention describes an isolated, purified or enriched nucleic
XX CC acid comprising a control region of a human peroxisome proliferator
XX CC activated receptor gamma (PPAR-gamma) gene. The nucleic acids are useful
XX CC for screening for agents capable of modulating the expression of a human
XX CC PPAR-gamma gene. These agents (modulators) form pharmaceutical
XX CC compositions that are useful for treating diseases associated with
XX CC high/low levels of human PPAR-gamma gene expression. The diseases include
XX CC obesity, anorexia, cachexia, lipodystrophy, lipomas, liposarcomas,
XX CC abnormalities associated with anti-human immunodeficiency virus (HIV)
XX CC treatment, insulin resistance, non-insulin-dependent diabetes mellitus
XX CC (NIDDM), polycystic ovary syndrome, diseases of the gastrointestinal (GI)
XX CC tract, inflammatory bowel disease, Crohn's disease, ulcerative colitis
XX CC and bowel cancer. The nucleic acids are useful for studying the role of
XX CC the PPAR-gamma gene in various diseases and disorders. The structure of
XX CC PPAR-gamma enables genetic studies of PPAR- gamma mutations in humans,
XX CC and evaluation of its role in disorders like insulin resistance, NIDDM,
XX CC and diseases associated with altered adipose tissue function, like
XX CC obesity and lipodystrophic syndromes. The nucleic acids are also useful
XX CC for gene therapy and the production of transgenic animals, which are
XX CC useful in screening assays. The control regions of the nucleic acids
XX CC enable screening for modulators of the human PPAR-gamma gene, which are
XX CC useful in designing drugs for treating disorders or diseases associated
XX CC with the level of PPAR-gamma gene expression. The present sequence
XX CC represents the human PPAR-gamma-2 proximal promoter
XX XX
XX SQ Sequence 177 BP; 44 A; 35 C; 29 G; 69 T; 0 U; 0 Other;

Query Match 10.3%; Score 52; DB 2; Length 177;
Best Local Similarity 100.0%; Pred. No. 0.00026;
Matches 52; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 452 GTCCTTCTGTGTTATCCCATCTCTCCCAATATTGGAACTGATGCT 503
|||||
Db 1 GTCCTTCTGTGTTATCCCATCTCTCCCAATATTGGAACTGATGCT 52
|||||

RESULT 8
ADO47191
ID ADO47191 standard; DNA; 200000 BP.
XX AC ADO47191;
XX AC
XX AC
XX DT 15-JUL-2004 (first entry)
XX XX
XX DE DNA sequence of a human immunoglobulin heavy chain variable region.
XX KW immunoglobulin; B cell; germline; region V; region D; region J;
XX KW recombination-promoting protein; recombination activating gene 1; RAG-1;
XX KW RAG-2; ss; terminal deoxynucleotidyltransferase; Tdt; V(D)J recombinase.
XX OS Homo sapiens.
XX XX

PN WO2004029249-A1.
XX PD 08-APR-2004.
XX PF 30-SEP-2003; 2003WO-AU001286.
XX PR 30-SEP-2002; 2002US-0415024P.
XX PA (CENT-) CENTENARY INST CANCER MEDICINE & CELL BI.
XX PI Jolly C;
XX DR WPI; 2004-316126/29.
XX XX
XX PT New vector comprising one or more immunoglobulin regions selected from V,
XX PT D, and J regions, a 5' flanking region, and a 3' flanking region, useful
XX PT for reverting cell lines to a pro-B cell-like state or to a germline-like
XX PT state.
XX XX
XX PS Example 1; Page 136-185; 382pp; English.
XX XX
XX CC The specification describes a method for the preparation of
XX CC immunoglobulins. The method uses a vector for reverting cell lines to a
XX CC pro-B cell-like state or to a germline-like state, by replacing one or
XX CC more of the immunoglobulin regions V, D, and J of the rearranged
XX CC immunoglobulin gene with the germ-line or unrearranged V, D or J regions.
XX CC The vector can comprise a polynucleotide sequence encoding a
XX CC recombination-promoting protein, selected from recombination activating
XX CC gene 1 (RAG-1), RAG-2, terminal deoxynucleotidyltransferase (tdt). These
XX CC proteins collectively constitute a V(D)J recombinase. The method is
XX CC useful for the preparation of immunoglobulin heavy and/or light chains
XX CC and their fragments. The present sequence encodes a human immunoglobulin
XX CC heavy chain variable region, and is used as a template to construct
XX CC vectors for use in the method of the invention.
XX XX
XX SQ Sequence 200000 BP; 59904 A; 41194 C; 41535 G; 57367 T; 0 U; 0 Other;

Query Match 9.0%; Score 45.2; DB 12; Length 200000;
Best Local Similarity 48.1%; Pred. No. 0.19;
Matches 128; Conservative 0; Mismatches 138; Indels 0; Gaps 0;

Qy 30 TCAGTAGCATGCTGATACCAACGTTTAAACATGATGATACATATTTGAATTCCAAATTTT 89
|||||
Db 139429 TGAGAACATGGAAAGCTTTCTTCTAAGATAAGGAATAAGAAAGATGCCAGTCTTA 139488
|||||
Qy 90 CTTGAGATAATGCTGATTAGAGATTAGAGATTCACAGGGATAGACACCGAAGAAACT 149
|||||
Db 139489 ATAATTCATACGATATTGATGATAGATTTCACCATAGAAATTTGAGAAAGAAATAA 139548
|||||
Qy 150 TTGCCCAATAAGCTTTCTCGTATTTCATAGCAAGAGATTTTAAGTTTTCATTTAAGAA 209
|||||
Db 139549 TAGTAAAGGCATACAAATTTGAGGGGAATAAGTTAAATTTCTCTGTTTACAGACAATAT 139608
|||||
Qy 210 GCATTTGTGAATTATCAACAATAAATAATGCAAGTGGAATTTGAACAGCTCTTCTCTG 269
|||||
Db 139609 AACTTTATAAATTCCAAAAAATAAATAATTTCTGAAAAAAGCAGCCAGTAAACAA 139668
|||||
Qy 270 ATAATTCATAATACAGTACAGTTTCAC 295
|||||
Db 139669 ACAATTTAGTAACATTCGAGTATAC 139694
|||||

RESULT 9
ABQ72637/c
ID ABQ72637 standard; CDNA; 3162 BP.
XX AC ABQ72637;
XX AC
XX DT 03-SEP-2002 (first entry)
XX XX
XX DE Human MDDT encoding cDNA SEQ ID NO 189.
XX OS Human; MDDT; disease detection and treatment molecule polynucleotide;
XX KW

KW proliferative disorder; hepatitis; psoriasis; cancer; AIDS;
KW autoimmune disorder; inflammatory disorder; allergy; multiple sclerosis;
KW rheumatoid arthritis; transgenic; gene therapy; antiarteriosclerotic;
KW hepatotropic; antiinflammatory; antipsoriatic; cytoskeletal; anti-HIV;
KW anti-allergic; antianaemic; antiaesthetic; antiatherosclerotic; antigout;
KW neuroprotective; antirheumatic; antiarthritic; gene; ss.
XX
OS Homo sapiens.
XX
XX WO200240715-A2.
XX
XX 23-MAY-2002.
XX
XX
XX 06-SEP-2001; 2001WO-US027628.
XX
XX 05-SEP-2000; 2000US-0229747P.
XX
XX 05-SEP-2000; 2000US-0229748P.
XX
XX 05-SEP-2000; 2000US-0229749P.
XX
XX 05-SEP-2000; 2000US-0229750P.
XX
XX 05-SEP-2000; 2000US-0229751P.
XX
XX 06-SEP-2000; 2000US-0230583P.
XX
XX 06-SEP-2000; 2000US-0230585P.
XX
XX 06-SEP-2000; 2000US-0230514P.
XX
XX 06-SEP-2000; 2000US-0230515P.
XX
XX 06-SEP-2000; 2000US-0230517P.
XX
XX 06-SEP-2000; 2000US-0230518P.
XX
XX 06-SEP-2000; 2000US-0230519P.
XX
XX 06-SEP-2000; 2000US-0230595P.
XX
XX 06-SEP-2000; 2000US-0230597P.
XX
XX 06-SEP-2000; 2000US-0230598P.
XX
XX 06-SEP-2000; 2000US-0230599P.
XX
XX 06-SEP-2000; 2000US-0230610P.
XX
XX 06-SEP-2000; 2000US-0230865P.
XX
XX 06-SEP-2000; 2000US-0230988P.
XX
XX 06-SEP-2000; 2000US-0230989P.
XX
XX 07-SEP-2000; 2000US-0230951P.
XX
XX 07-SEP-2000; 2000US-0231163P.
XX
XX 07-SEP-2000; 2000US-0231167P.
XX
XX (INCY-) INCYTE GENOMICS INC.
XX
XX Jackson S, Lincoln SE, Altus CM, Dufour GE, Chalup MS;
PI Hillman JL, Jones AU, Yu JY, Wright RU, Gietzen D, Liu TP, Yap PE;
PI Dahl CR, Momiya MG, Bradley DL, Rohatgi SD, Harris B;
PI Roseberry AM, Gerstein EH, Peralta CH, David MH, Panzer SR, Flores V;
PI Daffo A, Marwaha R, Chen AJ, Chang SC, Au AP, Inman RR;
XX
XX WPI; 2002-527544/56.
DR P-PSDB; ABP51420.
XX
XX Novel human disease detection and treatment polypeptide, useful in
PT diagnosis, prevention or treatment of cell proliferative disorders e.g.
PT arteriosclerosis, cirrhosis and an autoimmune/inflammatory disorder e.g.
PT AIDS.
XX
XX Claim 1; Page 407-408; 618pp; English.
XX
XX The invention relates to an isolated human disease detection and
CC treatment (MDPT) polypeptide (I) selected from a polypeptide having a
CC sequence selected from 254 sequences (ABP51231-ABP51484) given in the
CC specification, a naturally occurring polypeptide comprising a sequence
CC having at least 90% identity to (I) or a biologically active or
CC immunogenic fragment of (I). (I) is useful for screening a compound for
CC effectiveness as an agonist or antagonist, for screening a compound that
CC specifically binds (I) or modulates the activity of (I), and for
CC preparing a polyclonal or monoclonal antibody by hybridoma technology.
CC Nucleic acids (II) (ABQ72449-ABQ72700) encoding (I) are useful for
CC screening a compound for effectiveness in altering expression of a target
CC polynucleotide comprising. Oligonucleotides and antibodies are useful for
CC detecting MDPT in a sample or for assessing toxicity of a test compound,
CC in a diagnostic test for a condition or a disease associated with the
CC expression of MDPT in a biological sample, for detecting (I) in a sample,
CC and for purifying (I) from a sample. A composition comprising (I), an

CC agonist or antagonist is useful for treating a disease or condition
CC associated with decreased or increased expression of functional MDPT. (I)
CC or (II) are useful for diagnosing, treating or preventing disorders
CC associated with aberrant expression of MDPT, where the disorders are
CC selected from a cell proliferative disorder such as arteriosclerosis,
CC cirrhosis, hepatitis, psoriasis, and cancer and an
CC autoimmune/inflammatory disorder such as AIDS, Addison's disease,
CC allergy, anaemia, asthma, atherosclerosis, gout, multiple sclerosis or
CC rheumatoid arthritis. (II) are useful for creating knockin humanised
CC animals or transgenic animals to model human diseases, in somatic or
CC germline gene therapy, to generate a transcript image of a tissue or cell
CC type, for detecting differences in the chromosomal location due to
CC translocation or inversion among normal, carrier or affected individuals
CC and as hybridisation probes for mapping naturally occurring genomic
CC sequences
XX
XX Sequence 3162 BP; 971 A; 580 C; 568 G; 943 T; 0 U; 0 Other;
SQ
Query Match 8.8%; Score 44.2; DB 6; Length 3162;
Best Local Similarity 46.3%; Pred. No. 0.092;
Matches 145; Conservative 0; Mismatches 168; Indels 0; Gaps 0;
Qy 13 CTTAGGATATACATTTTCAGTACGCTGATACCAACGTTTAACTATGATACATAT 72
Db 2842 CTTGAAATATAAATATCTATAAATTTCTATGATACATAGTTTAAAAATTTTGAATTTT 2783
Qy 73 TTGAATTCCTCAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATTCAACACGAGATA 132
Db 2782 TCTAAATATATAAGAAACTTAAAAAAGACAGAGAAATGCTTCCCTTAACATGTTTTC 2723
Qy 133 GACACCGAAAGAAACTTTGCCCCAAATAAGCTTTCTGGTATTTCATAGCAAGAGATTTA 192
Db 2722 CTATCAGAAATGAACCTTTTTCAGCTAATTCACCTTCTAATTAAGTAAACACAGAGCTTGA 2663
Qy 193 AGTTTTCATTTAAGAACCCATTGTGATTATACAAATAAATAAATCAAGTGTGATATT 252
Db 2662 AATCTAAGTTTAAAGAGATCAAAAATAGTATACAACTTAAANAATCATATGTTCTT 2603
Qy 253 GAACAGTCTCTCTCTGATAATTTCTAATAACAGTACAGTTCACGCCCTTCACGAGACACT 312
Db 2602 GGCCTCAGAAATTTTATTTTAAAGTTATACAGGATCAGATTATGAGACTGTGCAACCT 2543
Qy 313 GAACATGTGTCA 325
Db 2542 GGGTAATCTGTCA 2530
RESULT 10
ABL33191/c
ID ABL33191 standard; DNA; 6215 BP.
XX
XX ABL33191;
XX
XX 26-MAR-2002 (first entry)
XX
XX Human immune system associated gene SEQ ID NO: 1164.
XX
XX Human; immune system disease; cytosine methylation; antiaesthetic;
KW antiarteriosclerotic; antianaemic; cytoskeletal; neutropenic;
KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;
KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;
KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;
KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;
KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;
ds.
XX
XX Homo sapiens.
OS
XX WO200200928-A2.
XX
XX 03-JAN-2002.
XX
XX 02-JUL-2001; 2001WO-EP007537.

```
XX 30-JUN-2000; 2000DE-01032529.
PR 01-SEP-2000; 2000DE-01043826.
XX (EPIG-) EPIGENOMICS AG.
XX Olek A, Piepenbrock C, Berlin K;
XX WPI; 2002-130909/17.
DR
XX Nucleic acid comprising fragment of chemically modified gene, useful for
PT diagnosis and treatment of diseases associated with abnormal cytosine
PT methylation.
XX
XX Claim 1; SEQ ID NO 1164; 32pp + Sequence Listing; German.
XX
XX The present invention provides a number of human immune system associated
CC genes which are modified by the methylation of cytosines. The sequences
CC can be used in the diagnosis and treatment of immune system disorders,
CC including eye diseases such as retinopathy, neovascular glaucoma and
CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
CC diseases. The present sequence is a gene of the invention
XX
XX SQ Sequence 6215 BP; 1813 A; 169 C; 1450 G; 2783 T; 0 U; 0 Other;
XX
XX Query Match 8.7%; Score 43.6; DB 6; Length 6215;
XX Best Local Similarity 58.5%; Pred. No. 0.17;
XX Matches 76; Conservative 0; Mismatches 54; Indels 0; Gaps 0;
XX
XX QY 167 CTGCTATTTCATAGCAAGACGATTTAAGTTTTCATTTAAGAACGCCATTGTGAATTATAC 226
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 5118 CTTTCATTTCTTACCATAAACCACATTTACCATTAAATTTAATATAAATACT 5059
XX
XX QY 227 AACATAAAATGCAAGTGATATTGAACAGTCTCTTCTCGATAATTTCTAAATACAGT 286
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 5058 AAAAACTCAAAAAAATAAACAACATTCCTCTCTTATACACATACATACTACAA 4959
XX
XX QY 287 ACAGTTTCAG 296
XX Db | |||||
XX 4998 ATTATTCAG 4989
XX
XX RESULT 11
XX ADI03931/c
XX ID ADI03931 standard; DNA; 99916 BP.
XX AC ADI03931;
XX DT 22-APR-2004 (first entry)
XX
XX DE Human enzyme protein encoding genomic DNA.
XX
XX KW Enzyme protein; drug screening; disease diagnosis; human; gene therapy;
XX chromosome 6; gene; ds; glucuronyltransferase.
XX
XX OS Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 2614..96915
XX FT /*tag= b
XX FT /product= "enzyme protein"
XX FT /note= "contains introns"
XX FT 2614..3204
XX FT /*tag= a
XX FT /number= 1
XX FT 3205..64312
XX FT /*tag= c
XX FT /number= 1
XX FT 64313..64457
XX FT /*tag= d
XX FT /number= 2
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FT intron 64458..96602
FT /*tag= e
FT /number= 2
FT exon 96603..96751
FT /*tag= f
FT /number= 3
FT intron 96752..96831
FT /*tag= g
FT /number= 3
FT exon 96832..96915
FT /*tag= h
FT /number= 4
XX WO200268657-A2.
XX
XX 06-SEP-2002.
XX
XX 08-FEB-2002; 2002WO-US003623.
XX
XX 26-FEB-2001; 2001US-0270871P.
XX 26-MAR-2001; 2001US-00816095.
XX (PEKE ) PE CORP NY.
XX
XX Gan W, Yan C, Merkulov GV, Di Francesco V, Beasley EM;
XX
XX WPI; 2002-713380/77.
XX P-PSDB; ADI03930.
XX
XX New human enzyme proteins, useful for treating or diagnosing disorders
XX associated with abnormal expression of the protein, in drug screening
XX assays and pharmacogenomic analysis.
XX
XX Claim 4; SEQ ID NO 3; 127pp; English.
XX
XX The invention relates to a novel isolated enzyme protein and encoding
XX polynucleotides. The protein shows a high degree of similarity to a
XX glucuronyltransferase cloned from a rabbit brain cDNA library. The
XX peptides and nucleic acid molecules are useful as models for the
XX development of human therapeutic targets, aid in the identification of
XX therapeutic proteins, and serve as targets for the development of human
XX therapeutic agents. The peptide may be used in drug screening assays, in
XX assays to determine the biological activity of the protein, to raise
XX antibodies or to elicit another immune response, as a reagent in assays
XX designed to quantitatively determine levels of the protein in biological
XX fluids, or as markers for tissues in which the corresponding protein is
XX preferentially expressed. The human enzyme protein is also useful for
XX diagnosing a disease, predisposition to a disease, or treating a disorder
XX characterized by an absence of, inappropriate or unwanted expression of
XX the protein. The antibodies are useful in pharmacogenomic analysis, for
XX inhibiting protein function, or for tissue typing. The nucleic acid
XX molecules are useful as probes, primers, chemical intermediates, or in
XX biological assays. The present sequence represents a human enzyme protein
XX encoding genomic DNA.
XX
XX SQ Sequence 99916 BP; 27736 A; 18701 C; 19334 G; 29032 T; 0 U; 5113 Other;
XX
```

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Query Match 8.3%; Score 41.8; DB 6; Length 99916;
Best Local Similarity 50.8%; Pred. No. 1.3;
Matches 100; Conservative 0; Mismatches 97; Indels 0; Gaps 0;
XX
XX QY 42 TGATACCAAGCTTTAAACTATGGATACATATTTGAATTCCAAATTTTCTTCAGATAATG 101
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 8890 TGTAACCTAACCTGCACATTGTGCACATGTACCCCTAAACCTTAAGTATATAATAATAA 8831
XX
XX QY 102 TGATTAGAGATTAGAGATTCAACAGGGATAGACACCGAAGAAANAACCTTGCCCAATAAA 161
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 8830 ATAAATAAAATAAAGAAATATCCAGGAGATCAACCCACAGACAAAGAAGCTAATAA 8771
XX
XX QY 162 GCTTCTCGGTATTTTCATACGAGAGATTTAAGTTTTCATTTAAGAGCGCATTTGCGAAT 221
XX Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 8770 AGTACATTTTCTTGCAAAAAAATAATTTTATTTTCAATGTATATACATTAACAACAA 8711
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Qy 194 GTTTCCTTAAAGCCATTGTGAATTATACAACTAAATAAAATGCAA 243
ADJ40655/c
ID ADJ40655 standard; cDNA; 2000 BP.
XX
XX
AC ADJ40655;
XX
XX 06-MAY-2004 (first entry)
XX
XX Plant cDNA #1655.
XX
XX Plant; gene; ss; transcription; plant genome augmentation; cereal;
KW soybean; alfalfa; sunflower; canola; cotton; peanut; tobacco; sugar beet;
KW maize; barley; sorghum; rice; wheat; crop plant; insecticide resistance;
KW stress tolerance; salt tolerance; cold tolerance; drought tolerance;
KW plant nutrition; apical dominance; dwarfism; early flowering; antiviral;
KW antifungal.
XX
OS Eukaryota.
XX
XX US2004016025-A1.
XX
XX 22-JAN-2004.
XX
XX 26-SEP-2002; 2002US-00260238.
XX
XX 26-SEP-2001; 2001US-0325277P.
XX 26-SEP-2001; 2001US-0325448P.
XX 04-APR-2002; 2002US-0370620P.
XX
XX (BUDW/) BUDWORTH P.
XX (MOUG/) MOUGHAMER T.
XX (BRIG/) BRIGGS S P.
XX (COOP/) COOPER B.
XX (GLAZ/) GLAZEBROOK J.
XX (GOFF/) GOFF S A.
XX (KATA/) KATAGIRI F.
XX (KREP/) KREPS J.
XX (PROV/) PROVART N.
XX (RICK/) RICHE D.
XX (ZHUT/) ZHU T.
XX
XX Budworth P, Moughamer T, Briggs SP, Cooper B, Glazebrook J;
XX Goff SA, Katagiri P, Kreps J, Provart N, Ricke D, Zhu T;
XX WPI; 2004-190374/18.
XX
XX New rice promoter, useful for manipulating crop plants to alter or
XX improve phenotypic characteristics, e.g. produce large quantities of oil
XX or proteins, resistance to insecticides, virus or fungi, stress tolerance
XX or high nutritional value.
XX
XX Claim 26; SEQ ID NO 1655; 230pp; English.
XX
XX The invention relates to plant nucleotide sequences that direct seed-,
XX leaf- and/or stem-, panicle-, root- or pollen-specific or -preferential
XX or constitutive transcription of an operatively linked nucleic acid
XX segment. The invention also relates to a method for augmenting a plant
XX genome and a method of identifying a gene, where its expression is
XX altered in the seed, leaf, stem, panicle, pollen, root or is constitutive
XX in a plant cell. The plant is a cereal, e.g. soybean, alfalfa, sunflower,
XX canola, cotton, peanut, tobacco or sugar beet, preferably maize, barley,
XX sorghum, rice or wheat. The polynucleotides and the polypeptides they
XX encode are useful for manipulating crop plants to alter or improve
XX phenotypic characteristics, to produce large quantities of oil or
XX proteins, to incur resistance to insecticides, viruses or fungi, and to
XX incur stress tolerance (e.g. salt, cold or drought) to ensure the plants
XX have a high nutritional value with reduced apical dominance or dwarfism,
XX early flowering or altered metabolic pathways. This sequence represents a

CC plant nucleic acid of the invention. Note: The sequence data for this
CC patent did not form part of the printed specification but was obtained in
CC electronic format directly from USPTO at seqdata.uspto.gov/sequence.html.
XX
SQ Sequence 2000 BP; 540 A; 381 C; 366 G; 713 T; 0 U; 0 Other;
Query Match 8.1%; Score 40.8; DB 12; Length 2000;
Best Local Similarity 48.0%; Pred. No. 0.69;
Matches 117; Conservative 0; Mismatches 127; Indels 0; Gaps 0;
Qy 4 AGTAAGTACCTTAGGAATATAACATTTTCAGTAGATGCTGATACCAACGTTTAAACTATG 63
Db 1937 ATTCGTATGATAAACATAGAAAATAATAAGATGAAGATGTGTTCTTTCAACTAAAT 1878
Qy 64 GATACATATTTGAATCCAAATTTTCTTCAGATAATGCTGATTAGAGATTAGATTCAA 123
Db 1877 GACACGATTTTGTATGCTTCATATATTTTATATCATCGAGAAAATCTATAAG 1818
Qy 124 CCAGGGATAGACACCCGAAAGAAAACCTTGCCCAAATAAGCTTTCTGGTATTTTCATAAGCA 183
Db 1817 CAATTGAGATGATTGCTTTATGATATTTTAGATCAATATTTGCTTCATCGCTAAAGGA 1758
Qy 184 AGAGATTTAAGTTTTCATTTAAGAACCCATTGTGAATTATACAACTAAATAATGCAA 243
Db 1757 AAAAAGCTAAGATGATGGGACACAAAGGACAATGTGGACCATAGAGCTATAAATAATTAAA 1698
Qy 244 GTGG 247
Db 1697 ATGG 1694
RESULT 15
ADQ97818/c
ID ADQ97818 standard; DNA; 109661 BP.
XX
XX ADQ97818;
XX
XX 07-OCT-2004 (first entry)
XX
XX Human cancer associated sequence HD11-002, SEQ ID 795.
XX
XX Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Human; ds.
XX
XX Homo sapiens.
XX
XX W02004060304-A2.
XX
XX 22-JUL-2004.
XX
XX 22-DEC-2003; 2003WO-US041389.
XX
XX 27-DEC-2002; 2002US-00330773.
XX
XX (SAGR-) SAGRES DISCOVERY INC.
XX
XX Morris DW, Malandro MS;
XX
XX WPI; 2004-543781/52.
XX
XX New isolated cancer associated nucleic acids comprising at least 10
XX contiguous nucleotides, useful for diagnosing, preventing and/or treating
XX cancers such as leukemia and lymphoma.
XX
XX Claim 1; SEQ ID NO 795; 199pp; English.
XX
XX The present invention relates to cancer associated sequences (ADQ97025-
XX ADQ98004). The sequences are useful for the diagnosis, prevention and/or
XX treatment of cancer, such as leukemia and lymphoma. Note: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 109661 BP; 30680 A; 19371 C; 19986 G; 35350 T; 0 U; 4274 Other;
XX

Search completed: November 9, 2005, 00:12:39
Job time : 411.412 secs

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Qy	360	TGTACCAAGTCTTGCCAGAGCAGTGAACAATTATGACAACAATTTTTGTGCACAGCTGGCTC	419
Db	592	TGTACCAAGTCTTGCCAAAGCAGTGAACATTATGACAACAATTTTTGTGCACAGCTGGCTC	651
Qy	420	CTAATAGGACAGTGCAGCCAAATTCAGGCCAGTCCTTTCTGTGTTATTTCCCATCTCTC	479
Db	652	CTAATAGGACAGTGCAGCCAAATTCAGGCCAGTCCTTTCTGTGTTATTTCCCATCTCTC	711
Qy	480	CCAAATATTTGGAACTGATGCT 503	
Db	712	CCAAATATTTGGAACTGATGCT 735	
RESULT 2			
US-09-949-016-15491			
; Sequence 15491, Application US/09949016			
; Patent No. 6812339			
; GENERAL INFORMATION:			
; APPLICANT: VENTER, J. Craig et al.			
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED			
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF			
; FILE REFERENCE: CL001307			
; CURRENT APPLICATION NUMBER: US/09/949,016			
; CURRENT FILING DATE: 2000-04-14			
; PRIOR APPLICATION NUMBER: 60/241,755			
; PRIOR FILING DATE: 2000-10-20			
; PRIOR APPLICATION NUMBER: 60/237,768			
; PRIOR FILING DATE: 2000-10-03			
; PRIOR APPLICATION NUMBER: 60/231,498			
; PRIOR FILING DATE: 2000-09-08			
; NUMBER OF SEQ ID NOS: 207012			
; SOFTWARE: FastSeq for Windows Version 4.0			
; SEQ ID NO 15491			
; LENGTH: 86877			
; TYPE: DNA			
; ORGANISM: Human			
US-09-949-016-15491			
Query Match 90.5%; Score 455.2; DB 4; Length 86877;			
Best Local Similarity 96.8%; Pred No. 1.1e-123;			
Matches 488; Conservative 0; Mismatches 8; Indels 8; Gaps 2;			
Qy	1	TAGAGTAACTACCTTAGGAATATAACAATTCAGTAGCATGCTGATACCAACGTTTAAACT	60
Db	1472	TAGAGTAACTACCTTAGGAATATAACAATTCAGTAGCATGCTGATACCAACGTTTAAACT	1531
Qy	61	ATGATACATATTTGAATTCCAATTTTTCTTCAGATATGTTAGATTTAGAGATT	120
Db	1532	ATGATACATATTTGAATTCCAATTTTTCTTCAGATATGTTAGATTTAGAGATT	1584
Qy	121	CAACCCAGGGATAGACACCGAAGAAAACTTTGCCCAAAATAGCTTTCTGGTATTTCAATA	180
Db	1585	CAACCCAGGAATAGACACCGAAGAAAACTTTGCCCAAAATAGCTTTCTGGTATTTCAATA	1644
Qy	181	GCAAGAGATTTAAGTTTTCCATTTTAAAGAGCCATTGTGAATTATACAAATAAAAAATG	240
Db	1645	GCAAGAGATTTAAGTTTTCCATTTTAAAGAGCAATTTGTGAATTTTCAATAAAAAATG	1704
Qy	241	CAAGTGGATATTTGAACAGCTCTCTCTCTGATAATTTCTAAATACAGTACAGTTTCACGCCCC	300
Db	1705	CAAGTGGATATTTGAACAGCTCTCTCTCTGATAATTTCTAAATACAGTACAGTTTCACGCCCC	1764
Qy	301	TCACGAGACACTGAAACATGT--GGTCAACGGCGAGACAGTGGGCAATATTTATCCCTGTAA	359
Db	1765	TCACGAGACACTGAAACATGTGGGTCAACGGCGAGACAGTGGGCAATATTTTCCCTGTAA	1824
Qy	360	TGTACCAAGTCTTGCCAGAGCAGTGAACATTATGACAACAATTTTTGTGCACAGCTGGCTC	419
Db	1825	TGTACCAAGTCTTGCCAAAGCAGTGAACATTATGACAACAATTTTTGTGCACAGCTGGCTC	1884
Qy	420	CTAATAGGACAGTGCAGCCAAATTCAGGCCAGTCCTTTCTGTGTTATTTCCCATCTCTC	479
Db	1885	CTAATAGGACAGTGCAGCCAAATTCAGGCCAGTCCTTTCTGTGTTATTTCCCATCTCTC	1944

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US-09-949-016-133111
; Sequence 133111, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 133111
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-133111

Query Match      28.3%; Score 142.4; DB 4; Length 601;
Best Local Similarity 99.3%; Pred. No. 3.5e-32;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 360 TGTACCAAGCTTCCAGAGCAGTGAACTATGACACAACTTTTGTGCAGCTGGCTC 419
Db 1 TGTACCAAGCTTCCAAAGCAGTGAACTATGACACAACTTTTGTGCAGCTGGCTC 60

Qy 420 CTAATAGGACAGTGCCAGCCCAATTCAGCCAGCTCTTCTGTGTATTCCCATCTCTC 479
Db 61 CTAATAGGACAGTGCCAGCCCAATTCAGCCAGCTCTTCTGTGTATTCCCATCTCTC 120

Qy 480 CCAAATATTTGGAACTGATGCT 503
Db 121 CCAAATATTTGGAACTGATGCT 144

RESULT 5
US-09-949-016-133194
; Sequence 133194, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 133194
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-133194

Query Match      28.3%; Score 142.4; DB 4; Length 601;
Best Local Similarity 99.3%; Pred. No. 3.5e-32;
Matches 143; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 360 TGTACCAAGCTTCCAGAGCAGTGAACTATGACACAACTTTTGTGCAGCTGGCTC 419
Db 1 TGTACCAAGCTTTCGCAAGCAGTGAACTATGACACAACTTTTGTGCAGCTGGCTC 60
```

```
Qy 420 CTAATAGGACAGTGCCAGCCCAATTCAGCCAGCTCTTCTGTGTATTCCCATCTCTC 479
Db 61 CTAATAGGACAGTGCCAGCCCAATTCAGCCAGCTCTTCTGTGTATTCCCATCTCTC 120

Qy 480 CCAAATATTTGGAACTGATGCT 503
Db 121 CCAAATATTTGGAACTGATGCT 144

RESULT 6
US-09-816-095-3/c
; Sequence 3, Application US/09816095
; Patent No. 6664084
; GENERAL INFORMATION:
; APPLICANT: GAN, Weiniu
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
; ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
; THEREOF
; FILE REFERENCE: CL001147
; CURRENT APPLICATION NUMBER: US/09/816,095
; CURRENT FILING DATE: 2001-03-26
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 99916
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(99916)
; OTHER INFORMATION: n = A,T,C or G
US-09-816-095-3

Query Match      8.3%; Score 41.8; DB 4; Length 99916;
Best Local Similarity 50.8%; Pred. No. 0.15;
Matches 100; Conservative 0; Mismatches 97; Indels 0; Gaps 0;

Qy 42 TGATACCAACGTTTAAACTATGGATACATATTTGCAATTTTCCAAATTTTCTTCAGATAATG 101
Db 8890 TGTAACCTAACCTGCACATTTGTGCACATGTACCTTAAAGCTTAAAGTATAATAATAA 8831

Qy 102 TGATTAGAGATTAGAGATTCAACAGGAGATAGACACCGAAGAAACTTTGCCCAATAA 161
Db 8830 ATAAATATAAATAAGAAATATCCAGGAAGATCAACCCACAGACAAAGAAAGCTAATAA 8771

Qy 162 GCTTCTGTGTTTTCATAAGCAAGAGATTAAAGTTTTCATTTTAAGAGCCATTGTGAAT 221
Db 8770 AGTACATTTCTTCGCAAAAAAATTTTATTTTCAATGTATATTAACAACAA 8711

Qy 222 TATACAAACAATAAAAAA 238
Db 8710 AAGGAAAAAGTAACAA 8694

RESULT 7
US-09-806-708B-22/c
; Sequence 22, Application US/09806708B
; Patent No. 6784342
; GENERAL INFORMATION:
; APPLICANT: The University of British Columbia
; TITLE OF INVENTION: Regulation of Embryonic Transcription in Plants
; FILE REFERENCE: 4810-58741
; CURRENT APPLICATION NUMBER: US/09/806,708B
; CURRENT FILING DATE: 2001-04-03
; PRIOR APPLICATION NUMBER: US 60/147,133
; PRIOR FILING DATE: 1999-08-04
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: Patent in version 3.0
; SEQ ID NO 22
; LENGTH: 1141
; TYPE: DNA
; ORGANISM: Artificial sequence
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; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16366
; LENGTH: 108310
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(108310)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-16366

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Query Match	7.6%;	Score 38.4;	DB 4;	Length 108310;
Best Local Similarity	47.2%;	Pred. No. 1.6;		
Matches 117;	Conservative 0;	Mismatches 131;	Indels 0;	Gaps 0;
Qy 42	TGATACCAACGTTTAAACTATG	GATACATATTTGAAATCCAAATTTTCTTC	CAGATAATG	101
Db 16475	TTATATATATTTATATATAAAATTTATG	TAAATTTATATATATATAAAATTTATAAA		16534
Qy 102	TGATTTAGAGNTTAGAGNTTCAAC	CAGGGATAGACCCGAAAGAAACTTTGCCCAAAATAA		161
Db 16535	TATATATAAANTTATATATATTTATATAAAATTTATAAAATTTATATAAAATTTAT			16594
Qy 162	GCTTCTCGGTATTTTCATAAGCAAGAGATTTAAAGTTTTC	CAATTTTAAAGAGCCCAATTTGTGAAT		221
Db 16595	AAATATATAAAAAATTTATAAATAATTTATAAATAATTTATAAATTTTATATAAAT			16654
Qy 222	TATACACAACTATAAAATGCAAGTGGATATTGACAGTCTCTTCT	CGATAATTCCTAAT		281
Db 16655	TATATATAAANTTATAAATATATATAAANTTATATAAATTTATATATATATAAATTTATAAATATATAT			16714
Qy 282	ACAGTACA			289
Db 16715	AAATTATA			16722

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RESULT 11
US-09-949-016-12756/c
; Sequence 12756, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12756
; LENGTH: 136264
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(136264)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-12756

```

Query Match 7.6%; Score 38.2; DB 4; Length 136264;
Best Local Similarity 49.8%; Pred. No. 2;

Matches	126;	Conservative	0;	Mismatches	123;	Indels	4;	Gaps	1;
Qy	66	TACATATTTGAATTC	CCAAATTTTCTT	CAGATAAATGTC	GATTAGAGATT	PAGAGNTTCA	ACC	125	
Db	24356	TATATAATATATAT	TAAATATATAT	TATATATATAT	TATATATATAT	TATATATAT	TATA	-- 24299	
Qy	126	AGGGATAGACACG	GAAGAAACTTTG	CCCAATAAGCTTT	CTGCTATTT	CATAAGCAAG	185		
Db	24298	--TAATATATAT	TATATATATAT	TATATATATAT	TATATATATAT	TATATATATAT	TATAT	24241	
Qy	186	AGATTTAAGTTTT	TCCATTTAAGA	CCCAATTGTGA	ATTATACAACA	ATAAAAAATG	CAAGT	245	
Db	24240	ATATATTAATAT	TATATATAT	TATATATAT	TATATATAT	TATATATAT	TATAT	24181	
Qy	246	GGATATTGAACAG	CTCTTCTGAT	TAATTTCTTA	ATATACAGT	TACAGTTAC	CGCCCTC	ACG 305	
Db	24180	TATATATATAT	TATATATAT	TATATATAT	TATATATAT	TATATATAT	TATAT	24121	
Qy	306	AGACACTGAACAT	318						
Db	24120	AGTAACCTCCACAT	24108						

RESULT 12

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US-09-949-016-13001/c
; Sequence 13001, Application US/09949016
; Patent NO. 6812339
;
GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
;
; FILE REFERENCE: CLO013307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
;
SEQ ID NO 13001
; LENGTH: 136265
; TYPE: DNA
; ORGANISM: Human
;
FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(136265)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13001

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Query Match	7.6%;	Score 38.2;	DB 4;	Length 136265;
Best Local Similarity	49.8%;	Pred. No. 2;		
Matches 126;	Conservative 0;	Mismatches 123;	Indels 4;	Gaps 1;
Qy	66	TACATATTTTGAATTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATTCACACC	125	
Db	24356	TATATAATATATATTAAATATATATATTATATAAATATATATTAAATTTATATATTATA	- 24299	
Qy	126	AGGGATAGACACGGAAGAAAACITTTGCCAAATAAGCTTTCTGTTATTTTCATAAGCAAG	185	
Db	24298	--TAAATATATATTAATAATATATATATATAATATAATATAATATAATATATATATATAT	24241	
Qy	186	AGATTTTAAGTTTTCCATTTTAAGAACCATTGTGAATTATACAACAATAAAAAATGCAAGT	245	
Db	24240	ATATATTAATATATATATATATATAATATAATATAATATAATATAATATAATATAATATAT	24181	
Qy	246	GGATATTGACAGTCTCTCTCTGATAATTTCTAAATACAGTACAGTTTCACGCCCTCAGG	305	
Db	24180	TAAATATATATATATATAAATAATACTATATATATATATATATATATATATATATATATAT	24121	
Qy	306	AGACACTGAAACAT	318	

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; Sequence 1010, Application US/09248796A
; Patent No. 6747137
; GENERAL INFORMATION:
; APPLICANT: Keith Weinstock et al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO CANDIDA ALBICAN
; FILE REFERENCE: 107196.132
; CURRENT APPLICATION NUMBER: US/09/248,796A
; PRIOR FILING DATE: 1999-02-12
; PRIOR APPLICATION NUMBER: US 60/074,725
; PRIOR FILING DATE: 1998-02-13
; PRIOR APPLICATION NUMBER: US 60/096,409
; PRIOR FILING DATE: 1998-08-13
; NUMBER OF SEQ ID NOS: 28208
; SEQ ID NO 1010
; LENGTH: 1317
; TYPE: DNA
; ORGANISM: Candida albicans
US-09-248-796A-1010
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Best Local Similarity 47.8%; Pred. No. 0.34;
Matches 110; Conservative 0; Mismatches 120; Indels 0; Gaps 0;
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Qy      113 TAGAGATTCACACAGGGATAGACACCGAAGAAACCTTTGCCAAATAAGCTTTCTGGTA 172
Db      684 TTTAGATATAAATTTAAATTTAGAACAAACAAATTTCCACTACCAAGAAATTTTAATGAA 743
Qy      173 TTTTCATAGCAGAGATTTAGTTTTCATTTTAAAGAGCCATTCGTAATTATACAAAT 232
Db      744 ATGGAGTATTAATTAATTAATGATTTTAAATTTACAAATACTAATGAATTTAAATTAAT 803
Qy      233 AAAAAATCGAAGTGATATTCGAACAGATCTCTTCTCTGATATTTCTAAATA 282
Db      804 TAATAACAATCGTGTGTAATCGGAATCATTTTTTAATTAATATGTTTATA 853
RESULT 14
US-09-790-988-1/c
; Sequence 1, Application US/09790988
; Patent No. 6632935
; GENERAL INFORMATION:
; APPLICANT: SHIGENOBU, SHUJI
; APPLICANT: WATANABE, HIDEMI
; APPLICANT: HATTORI, MASAHIRA
; APPLICANT: SAKAKI, YOSHIYUKI
; TITLE OF INVENTION: GENOME DNA OF BACTERIAL SYMBIONT OF APHIDS
; FILE REFERENCE: 081356/0159
; CURRENT APPLICATION NUMBER: US/09/790,988
; PRIOR FILING DATE: 2001-02-23
; PRIOR APPLICATION NUMBER: JP2000-107160
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 640681
; TYPE: DNA
; ORGANISM: Buchnera sp.
US-09-790-988-1
Query Match      7.5%; Score 37.6; DB 4; Length 640681;
Best Local Similarity 49.5%; Pred. No. 5.6;
Matches 97; Conservative 0; Mismatches 99; Indels 0; Gaps 0;
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Qy      104 ATTAGAGATTAGAGATTTCAACACGAGGATAGACACCGAAGAAACCTTTGCCAAATAAGC 163
Db      212203 AATAATGATCATTTATGTTATATAAAATTAATGCAATCTAAAAAGTTTTTAAAAAACAGTTC 212144
Qy      164 TTTCTGTTATTTCAATAGCAGAGATTTAAGTTTCCATTTTCCATTTAAGAACCCATTGTAATTA 223
Db      212143 TTTTAAATATTTTAAAAAATGAAAAATATATATATTTTATATAAAAAATATAATTTCTTTTA 212084
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; Sequence 15830, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15830
; LENGTH: 192506
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15830
Query Match      7.4%; Score 37; DB 4; Length 192506;
Best Local Similarity 52.2%; Pred. No. 5.1;
Matches 82; Conservative 0; Mismatches 75; Indels 0; Gaps 0;
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Db      191028 GGAAGAGACCATTTAACTCTCCAAACAATAGTGACATCAAGAAAAATGACACCAATTTG 190969
Qy      77 ATTCCAAATTTTCTTCCAGATAATGTGATTAGAGATTTCACACCGGGATAGACA 136
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SUMMARIES

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c 2	455.2	90.5	165043	19	US-10-235-192A-46
3	45.2	9.0	200000	20	US-10-672-764A-31
c 4	44.2	8.8	3162	20	US-10-363-829-189
c 5	43.6	8.7	6215	16	US-10-311-455-1164

Sequence 10, Appl

Sequence 46, Appl

Sequence 31, Appl

Sequence 189, App

Sequence 1164, App

c 6	41.8	8.3	99916	9	US-09-816-095-3	Sequence 3, Appli
c 7	41.8	8.3	99916	19	US-10-634-905-3	Sequence 3, Appli
c 8	41.2	8.2	32392	22	US-10-706-635-27	Sequence 27, Appl
c 9	40.8	8.1	2000	18	US-10-260-238-1655	Sequence 1655, Ap
c 10	40.8	8.1	59914	20	US-10-741-601-5619	Sequence 5619, Ap
c 11	40.8	8.1	101782	20	US-10-741-601-5661	Sequence 5661, Ap
c 12	40.4	8.0	1886	20	US-10-437-963-36653	Sequence 36653, A
c 13	40.2	8.0	797	19	US-10-424-599-18984	Sequence 18984, A
c 14	40	8.0	8056	21	US-10-473-126-240	Sequence 240, App
c 15	39.8	7.9	350	20	US-10-674-124A-1055	Sequence 1055, App
c 16	39.4	7.8	350	18	US-10-242-535A-53661	Sequence 53661, A
c 17	39.4	7.8	350	18	US-10-085-783A-53661	Sequence 53661, A
c 18	39.4	7.8	6191	16	US-10-311-455-1189	Sequence 1189, Ap
c 19	39.2	7.8	629	13	US-09-925-065A-861470	Sequence 861470, A
c 20	39.2	7.8	967	13	US-09-925-065A-37192	Sequence 37192, A
c 21	39.2	7.8	967	13	US-09-925-065A-37193	Sequence 37193, A
c 22	39.2	7.8	1943	13	US-09-925-065A-92097	Sequence 92097, A
c 23	39.2	7.8	6308	16	US-10-311-455-1443	Sequence 1443, Ap
c 24	39	7.8	6565	19	US-10-221-714A-188	Sequence 188, App
c 25	39	7.8	37515	20	US-10-433-793-28	Sequence 28, Appl
c 26	38.8	7.7	505	14	US-10-027-632-248390	Sequence 248390, A
c 27	38.8	7.7	505	18	US-10-027-632-248390	Sequence 248390, A
c 28	38.8	7.7	1696	18	US-10-374-780A-1530	Sequence 1530, Ap
c 29	38.8	7.7	1696	19	US-10-412-698B-1620	Sequence 1620, Ap
c 30	38.8	7.7	8056	21	US-10-473-126-386	Sequence 386, App
c 31	38.8	7.7	10945	16	US-10-240-453-228	Sequence 228, App
c 32	38.8	7.7	19634	21	US-10-473-126-156	Sequence 156, App
c 33	38.8	7.7	19634	21	US-10-473-126-302	Sequence 302, App
c 34	38.4	7.6	2258	26	US-11-097-143-29929	Sequence 29929, A
c 35	38.2	7.6	586	14	US-10-027-632-197296	Sequence 197296, A
c 36	38.2	7.6	586	18	US-10-027-632-197296	Sequence 197296, A
c 37	38.2	7.6	4152	22	US-10-706-635-9	Sequence 9, Appli
c 38	38.2	7.6	50000	19	US-10-706-635-25	Sequence 25, Appl
c 39	38.2	7.6	96589	22	US-10-052-482-214	Sequence 214, App
c 40	38	7.6	629	13	US-09-925-065A-861469	Sequence 861469, A
c 41	38	7.6	629	13	US-09-925-065A-897143	Sequence 897143, A
c 42	38	7.6	2035	24	US-10-795-159-412	Sequence 412, App
c 43	38	7.6	3426	13	US-09-925-065A-676913	Sequence 676913, A
c 44	38	7.6	3426	13	US-09-925-065A-676914	Sequence 676914, A
c 45	38	7.6	908765	24	US-10-795-159-685	Sequence 685, App

ALIGNMENTS

RESULT 1

US-10-181-176-10

; Sequence 10, Application US/10181176

; Publication No. US20050186567A1

; GENERAL INFORMATION:

; APPLICANT: Robert McKay

; APPLICANT: Alexander H. Borchers

; APPLICANT: Brenda F. Baker

; TITLE OF INVENTION: ANTISENSE MODULATION OF PEROXISOME PROLIFERATOR-ACTIVATED RECEPTOR

; TITLE OF INVENTION: GAMMA EXPRESSION

; FILE REFERENCE: RTSP-0328

; CURRENT FILING DATE: 2002-07-12

; PRIOR FILING DATE: 2001-01-11

; PRIOR APPLICATION NUMBER: US09/484,345

; PRIOR FILING DATE: 2000-01-18

; NUMBER OF SEQ ID NOS: 90

; SEQ ID NO 10

; LENGTH: 1100

; TYPE: DNA

; ORGANISM: Homo sapiens

; FEATURE:

; NAME/KEY: CDS

; LOCATION: (859)...(940)

US-10-181-176-10

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Qy 61 ATGGATACATATTTGAAATTTCCAAATTTTCTCTCAGATAATGCTGATTAGAGATTAGAGATT 120
Db 299 ATGGATACATATTTGAAATTTCCAAATTTTCTCTCAGATAATGCTGATTAGAGATT 351
Qy 121 CAACAGGGATAGACACCGAAGAAAACCTTTGCCAAATAAGCTTTCTGGTATTTCATAA 180
Db 352 CAACAGGGATAGACACCGAAGAAAACCTTTGCCAAATAAGCTTTCTGGTATTTCATAA 411
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Qy 241 CAAGTGGATATTGAACAGTCTCTCTCTGATAATTTCTAAATACAGTACAGTTACGCCCC 300
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Qy 301 TCACGAGACACTGAACATGT-GGTCAACCGCGAGACAGTGTGGCAATATTATCCCTGTAA 359
Db 532 TCACGAGACACTGAACATGTGGGTCAACCGCGAGACAGTGTGGCAATATTATCCCTGTAA 591
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Db 592 TGTACCAAGTCTTCCAAAGCAGTGAACATATGACACAACCTTTTGTCAAGTGGCTC 651
Qy 420 CTAATAGGACAGTGCAGCCCAATTTCAAGCCAGTCTCTCTGTTGTTTATCCCATCTCTC 479
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Qy 480 CCAAATATTGGAAACTGATGTCT 503
Db 712 CCAAATATTGGAAACTGATGTCT 735
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US-10-235-192A-46/c
; Sequence 46, Application US/10235192A
; Publication No. US20040043389A1
; GENERAL INFORMATION:
; APPLICANT: McCarthy, Jeanette
; TITLE OF INVENTION: Methods and Compositions for Identifying
; TITLE OF INVENTION: Risk Factors for Abnormal Lipid Levels and the Diseases
; TITLE OF INVENTION: and Disorders Associated Therewith
; FILE REFERENCE: MMI-011
; CURRENT APPLICATION NUMBER: US/10/235,192A
; CURRENT FILING DATE: 2002-09-04
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46
; LENGTH: 166043
; TYPE: DNA
; ORGANISM: Homo sapiens
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Best Local Similarity 96.8%; Pred. No. 1.3e-107;
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Qy 61 ATGGATACATATTTGAAATTTCCAAATTTTCTCTCAGATAATGCTGATTAGAGATTAGAGATT 120
Db 145729 ATGGATACATATTTGAAATTTCCAAATTTTCTCTCAGATAATGCTGATTAGAGATT 145677
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Db 145729 ATGGATACATATTTGAAATTTCCAAATTTTCTCTCAGATAATGCTGATTAGAGATT 145677
US-10-672-764A-31
; Sequence 31, Application US/10672764A
; Publication No. US20040156832A1
; GENERAL INFORMATION:
; APPLICANT: Jolly, Chris
; TITLE OF INVENTION: Immunoglobulin Compositions and Methods
; FILE REFERENCE: 13311.1001U
; CURRENT APPLICATION NUMBER: US/10/672,764A
; CURRENT FILING DATE: 2003-09-26
; NUMBER OF SEQ ID NOS: 68
; SOFTWARE: FastSeq for Windows Version 4.0
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; LENGTH: 200000
; TYPE: DNA
; ORGANISM: Human
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Best Local Similarity 48.1%; Pred. No. 2.6;
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Db 139429 TGAGAACATGGAAAGCTTTCTCTTAAGTAAAGATAGAAAGATGCCCAGTCTTA 139488
Qy 90 CTTTCAGATAATGTGATTAGAGATTAGAGATTCAACAGGAGATAGACACCGAAGAAAACCT 149
Db 139489 ATAATTTCTAATCAGCATATTGATGATGTTTAAACCATGAAATTTGAGAAAGAAAAA 139548
Qy 150 TTGCCCAATAAGCTTTCTGGTATTTCATAGCAAGAGATTAAAGTTTTCATTTAAGAA 209
Db 139549 TAGTAAAGGCATACAAAATTTGAGGGGAATTAAGTTAAATTTCTGTGTTACAGCAATAT 139608
Qy 210 GCCATTGTGAATTTATCAACAATAAAAAATGCAAGTGGATATTGAACAGTCTCTCTCTG 269
Db 139609 AACTTTATAAATTTCCAAAAAATAAAAAATTTCTGAAAAAACAGCCACTGAAACTAACAA 139668
Qy 270 ATAATTTCTAATAACAGTACAGTTTCAAC 295
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US-10-363-829-189/c
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QY	313	GAACATGTGGTCA	325
Db	2542	GGTAATCTGTCA	2530
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; Sequence 1164, Application US/10311455			
; Publication No. US20030143606A1			
; GENERAL INFORMATION:			
; APPLICANT: OLEK, Alexander			
; APPLICANT: PIEPENBROCK, Christian			
; APPLICANT: BERLIN, Kurt			
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System			
; TITLE OF INVENTION: cytosine methylation			
; FILE REFERENCE: 5013.1014			
; CURRENT APPLICATION NUMBER: US/10/311,455			
; CURRENT FILING DATE: 2002-12-16			
; PRIOR APPLICATION NUMBER: PCT/EP01/07537			
; PRIOR FILING DATE: 2001-07-02			
; PRIOR APPLICATION NUMBER: DE 10032529.7			
; PRIOR FILING DATE: 2000-06-30			
; PRIOR APPLICATION NUMBER: DE 10043826.1			
; PRIOR FILING DATE: 2000-09-01			
; NUMBER OF SEQ ID NOS: 2424			
; SEQ ID NO 1164			
; LENGTH: 6215			
; TYPE: DNA			
; ORGANISM: Artificial Sequence			
; FEATURE:			
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)			
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Best Local Similarity 58.5%; Pred. No. 1.5;			
Matches 76; Conservative 0; Mismatches 54; Indels 0; Gaps 0;			
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; Sequence 3, Application US/09816095
; Patent No. US20020137164A1
; GENERAL INFORMATION:
; APPLICANT: GAN, Weinhu
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
; TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001147
; CURRENT APPLICATION NUMBER: US/09/816,095
; CURRENT FILING DATE: 2001-03-26
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 99916
; TYPE: DNA
; ORGANISM: Human

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; OTHER INFORMATION: Clone ID:
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US-10-437-963-36653


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Db 208 CATGTAACATATATACACATATTTTATATATATATATATATATATATATATATTTTGTATATTT 149
Qy 111 ATTAGAGATTCAACCAGGATAGACACCGAAGAAACTTTGCCCAATAAGCTTCTGG 170
Db 148 ATTATATATATATATATATATATATATATATATATATATATATATATATATATATTTCTTATCCAG 89
Qy 171 TATTTTCATAAGCAAGAGATTTAAGTTTCCATTTAAGAAGCCATTGTGAATTATACAACA 230
Db 88 TCATTCATCTATAGATACTTAGGTGTTTTTATTTCTTGGCTGTGTGGAATAATGCTGCA 29
Qy 231 ATAAAAA 237
Db 28 ATGAATA 22
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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*

- 1: gb_est1.*
- 2: gb_est2.*
- 3: gb_hic.*
- 4: gb_est3.*
- 5: gb_est4.*
- 6: gb_est5.*
- 7: gb_est6.*
- 8: gb_gss1.*
- 9: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	51.6	10.3	1101	9	CNS00E47
C 2	48.2	9.6	1101	9	CNS0039G
C 3	46.4	9.2	747	8	BH957102
C 4	45.6	9.1	1106	9	CNS04LKI
C 5	44.6	8.9	766	8	BH460886
C 6	43.4	8.6	843	9	CNS00CS1
C 7	43.2	8.6	495	9	CC764640
C 8	43.2	8.6	618	9	CL335225
C 9	43.2	8.6	1201	9	CNS0167M
C 10	42.6	8.5	623	8	BH742342
C 11	42.4	8.4	669	8	BZ465736
C 12	42.4	8.4	818	8	BH661305
C 13	42.2	8.4	766	8	AQ544187
C 14	42.2	8.4	874	9	CG019400
C 15	42.2	8.4	1101	9	CNS000B8
C 16	42	8.3	473	6	CB888081
C 17	42	8.3	959	9	CNS0170R
C 18	42	8.3	1101	9	CNS006X3
C 19	41.8	8.3	266	9	CL883428
C 20	41.8	8.3	496	8	BH711021
C 21	41.8	8.3	591	8	BH467264
C 22	41.4	8.2	448	7	CK732164
C 23	41.4	8.2	617	9	CL625152
C 24	41.4	8.2	628	9	CL611954

C 25	41.2	8.2	1174	9	CL110860
C 26	41	8.2	724	7	CV464338
C 27	41	8.2	841	9	CR812616
C 28	41	8.2	1086	9	CNS00YXK
C 29	41	8.2	1123	8	CC273164
C 30	40.8	8.1	321	4	BG438771
C 31	40.8	8.1	412	5	BP623936
C 32	40.8	8.1	538	5	BH811789
C 33	40.8	8.1	693	9	CL934409
C 34	40.8	8.1	777	6	CB180244
C 35	40.8	8.1	832	8	BZ601497
C 36	40.8	8.1	1101	9	CNS00E34
C 37	40.8	8.1	1101	9	CNS00ZB7
C 38	40.6	8.1	705	8	BH970382
C 39	40.6	8.1	734	9	CNS010MP
C 40	40.6	8.1	736	8	BZ469037
C 41	40.6	8.1	1101	9	CNS00Z62
C 42	40.4	8.0	515	2	BE238266
C 43	40.4	8.0	738	6	CA104351
C 44	40.4	8.0	784	4	BI738064
C 45	40.2	8.0	326	4	BG370931

ALIGNMENTS

RESULT 1
LOCUS CNS00E47 1101 bp DNA linear GSS 04-JUN-1999
DEFINITION Drosophila melanogaster genome survey sequence TET3 end of BAC # BAC28M18 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION AL068720
VERSION AL068720.1 GI:4948863

KEYWORDS GSS.

SOURCE Drosophila melanogaster (fruit fly)

ORGANISM Drosophila melanogaster

Rukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;

Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;

Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 (bases 1 to 1101)

AUTHORS Genoscope.

TITLE Direct Submission

JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)

COMMENT - Web : www.genoscope.cns.fr
Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP).

The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osogawa and Aaron Mamoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2, a cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES
Location/Qualifiers

1..1101
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BAC28M18"
/note="end : TET3"

ORIGIN

Query Match 10.3%; Score 51.6; DB 9; Length 1101;
Best Local Similarity 29.8%; Pred. No. 0.011;


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Qy 130 ATAGACACCGAAGAACTTTGGCCCAATTAAGCTTTCTGTAATTTTCATAGCAAGAGAT 189
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 171 ATAAAAATTTAACTAACATAAAGGCACAAAAATAATTTTTTTTATAGTTGTACAT 230
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 190 TTAAGTTTTCAC 201
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 231 TTAATTTTACAA 242
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 4
CNS041KI
LOCUS CNS041KI 1106 bp DNA linear GSS 01-SEP-2000
DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
118P23 of library G from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL296235
VERSION AL296235.1 GI:9034815
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontoidea; Tetraodontidae; Tetraodon.
REFERENCE 1
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bertot,A., Fizeser,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Estimate of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
JOURNAL Nat. Genet. 25 (2), 235-238 (2000)
MEDLINE 20296633
PubMed 10835645
REFERENCE 2
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Ozouf-Costaz,C.,
Fizeser,C., Fischer,C., Bouneau,L., Billault,A., Quetier,F.,
Saurin,W., Bertot,A. and Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Genome Res. 10 (7), 939-949 (2000)
MEDLINE 20359837
PubMed 10899143
REFERENCE 3 (bases 1 to 1106)
GENOSCOPE
Direct Submission
Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
FEATURES
source
Location/Qualifiers
1..1106
/organism="Tetraodon nigroviridis"
/mol_type="genomic DNA"
/db_xref="taxon:99883"
/clone="118P23"
/clone_lib="G"
/note="Genoscope sequence ID : C08G118CH12LP1-end : T7"

ORIGIN
Query Match 9.1%; Score 45.6; DB 9; Length 1106;
Best Local Similarity 41.4%; Pred. No. 0.41;
Matches 116; Conservative 24; Mismatches 140; Indels 0; Gaps 0;

Qy 20 ATATAACATTTTCAGTAGCATGCTGATACCAAGTTTAAACATGATGATACATATTGTAAT 79
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 777 AAAAAAAATTTTCTTTTCTTTTAAACCAACTTACACACAAATTTTCTTTTAA 836
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 80 CCATATTTTCTTCAGTAATGTGATTAGAGATTAGAGATTCAACCGGATAGACCG 139
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

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Db 837 ACAAAWAATTTTTCACAAATTTWACAAAAAAATTTWTATCATCAATAAAAAACAACCT 896
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 140 AAAGAAAATTTTGGCCCAATTAAGCTTTCTGTAATTTTCATAGCAAGAGATTTAAGTTTC 199
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 897 YTTAAAAACTTTTAAAAAHYTTTATWTWACCTTTTAACTTTTAAAAAACYTTT 956
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 200 CATTTAAGAGCCATTTGTGAATTTATACAAATAAAAATGCAAGTGGATATTGAACAGT 259
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 957 TAAAAACACCACTCTCCYACAAACWCACAAATAAAAAAAWTAATTAATTAATTTT 1016
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 260 CTCTTCTCTGATTAATTTAAATACAGTACAGTTCACGCC 299
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 1017 TTTBTTTTTTTTTTAATTTTATTWAAAWAAACAAACACCC 1056
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 5
BH460886/C
LOCUS BH460886 766 bp DNA linear GSS 13-DEC-2001
DEFINITION BOGSK22TF BOGS Brassica oleracea genomic clone BOGSK22, genomic
survey sequence.
ACCESSION BH460886
VERSION BH460886.1 GI:17650631
KEYWORDS GSS.
SOURCE Brassica oleracea
ORGANISM Brassica oleracea
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Brassica.
REFERENCE 1 (bases 1 to 766)
AUTHORS Town,C.D., Van Aken,S., Utterback,T., Koo,H. and Fraser,C.M.
TITLE Whole genome shotgun sequencing of Brassica oleracea
JOURNAL Unpublished (2001)
COMMENT Other GSSs: BOGSK22TR
Contact: Chris Town
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.
Tel: 301-838-3523
Fax: 301-838-0208
Email: cdtown@tigr.org
DNA is from a doubled haploid provided by Tom Osborn.
Seq primer: TF
Class: sheared ends.
FEATURES
source
Location/Qualifiers
1..766
/organism="Brassica oleracea"
/mol_type="genomic DNA"
/strain="T01000DH3"
/db_xref="taxon:3712"
/clone="BOGSK22"
/clone_lib="BOGS"
/note="vector: PHOS1, Site 1: BstXI; 2-3 kb sheared
genomic DNA inserted into PHOS1 using BstXI linkers"

ORIGIN
Query Match 8.9%; Score 44.6; DB 8; Length 766;
Best Local Similarity 49.4%; Pred. No. 0.7;
Matches 116; Conservative 0; Mismatches 119; Indels 0; Gaps 0;

Qy 27 ATTTTCAGTAGCATGCTGATACCAAGTTTAAACTATGATGATACATATTGTAATTC 86
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 708 ATAACATATATTTTGTATATAATTTTATTAAGTGTATTATTAATGTAATTAATACTATA 649
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 87 TTTCTTCAGATAATGTGATTAGAGATTAGAGATTCAACCGGATAGACACCGAAGAAA 146
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 648 ATATATAACATATTTCTGTTAGATATAACATATGTAACTTGTAAAAATTAATAAAAA 589
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 147 ACTTTGCCCAATTAAGCTTTCTGTAATTTTCATAGCAAGAGATTTAAGTTTCCATTTAA 206
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 588 CAATGCCGAAATATTTTGTCTCAAAATTTTCTTTGTTTATTTATATATACATATTT 529
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 207 GAAGCCATTGTGAATTTATACAACTAAATAAATGCAAGTGGATTTGAACAGTCT 261
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 528 TAGGAATAATATACATAAACTATGAGCAACAACTAATTTGATTATTGAACTTTT 474
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

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RESULT 6	CNS00CS1	843 bp	DNA	linear	GSS 04-JUN-1999
LOCUS	Drosophila melanogaster genome survey sequence TET3 end of BAC #				
DEFINITION	BACR26H19 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.				
ACCESSION	AL059666				
VERSION	AL059666.1	GI:4947129			
KEYWORDS	GSS.				
SOURCE	Drosophila melanogaster (fruit fly)				
ORGANISM	Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.				
REFERENCE	1 (bases 1 to 843)				
AUTHORS	Genoscope.				
TITLE	Direct Submission				
JOURNAL	Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr - Web : www.genoscope.cns.fr)				
COMMENT	Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see http://www.fruitfly.org The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osoegawa and Aaron Mammoler in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm .				
FEATURES	Location/Qualifiers				
source	1..843				
	/organism="Drosophila melanogaster"				
	/mol_type="genomic DNA"				
	/db_xref="taxon:7227"				
	/clone="BACR26H19"				
	/clone_lib="RPCI-98"				
	/note="end : TET3"				
ORIGIN					
Query Match	8.6%;	Score 43.4;	DB 9;	Length 843;	
Best Local Similarity	36.7%;	Pred. No. 1.5;			
Matches	90;	Conservative	39;	Mismatches	116; Indels 0; Gaps 0;
QY	38	ATGCTGATACCAACGTTTAAACTATGGATACATATTTGAAATTTCCAAATTTTCTTCAGAT	97		
DB	578	WTTTATATATATATTTTWWTTTTTTTAYAWAAATTTAATTTATATATTTATATAT	637		
QY	98	AATGATGATTAGAGATTCAACAGGGATAGACACCGAAGAAACCTTTGCCCAA	157		
DB	638	WTTATWAAWAAWATATWAAWATAAAWATAAATATWTTWAAATAATTAATTAATW	697		
QY	158	ATAAGCTTCTCGTATTTTCATAAGCAAGAGATTTTAAAGTTTTCATTTTAAAGACCCAT	217		
DB	698	TATATATATATATATATATATWAAATTTWAAWATATATATATATATATATATATAT	757		
QY	218	GAATTTATACAAATAAAAAATGCAAGTGGATATTTGAACACAGTCTCTTCTCGATAAT	277		
DB	758	AAAAAAWATWATAAATAAAWAAATWATAYAWTTATTHWAAWAAATAATTAATAAA	817		
QY	278	AAATA	282		
DB	818	ATATW	822		
RESULT 7					
LOCUS	CC764640	495 bp	DNA	linear	GSS 27-JUN-2003
DEFINITION	CH240_47L12.TJ CHORI-240 Bos taurus genomic clone CH240_47L12, genomic survey sequence.				
ACCESSION	CC764640				
VERSION	CC764640.1	GI:32311143			
KEYWORDS	GSS.				
SOURCE	Bos taurus (cow)				
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.				
REFERENCE	1 (bases 1 to 495)				
AUTHORS	Larkin,D.M., Everts-van der Wind,A., Rebeiz,M., Schweitzer,P., Bachman,S., Green,S., Campos,E.J., Benson,L.D., Edwards,J., Liu,L., Womack,J.E., de Jong,P.J. and Lewin,H.A.				
TITLE	Bovine BAC end sequences from CHORI-240 library				
JOURNAL	Unpublished (2003)				
COMMENT	Other GSSs: CH240_47L12.TV Contact: Harris Lewin Department of Animal Sciences University of Illinois at Urbana Champaign 1201 W. Gregory Dr., Urbana, IL 61801, USA Tel: 217 333 5998 Fax: 217 244 5617 Email: h-lewin@uiuc.edu Clones are derived from the bovine BAC library CHORI-240 (http://www.chori.org/bacpac/bovine240.htm). For BAC library availability, please contact Pieter de Jong (pdejong@mail.cho.org). Clones may be purchased from BACPAC Resources (http://www.chori.org/bacpac/ordering/information.htm). This work was undertaken as part of the International Bovine BAC Mapping Consortium (IBBMC) by University of Illinois at Urbana Champaign, USA with funds provided by grant No. AG202-34480-11828 from USDA-CSREES and AG99-35205-8534 from USDA/NRI (Livestock Genome Sequencing Initiative) Plate: 47 row: L column: 12 Seq primer: SP6 Class: BAC ends.				
FEATURES	Location/Qualifiers				
source	1..495				
	/organism="Bos taurus"				
	/mol_type="genomic DNA"				
	/strain="breed: Hereford"				
	/db_xref="taxon:9913"				
	/clone="CH240_47L12"				
	/sex="Male"				
	/cell_type="Blood"				
	/clone_lib="CHORI-240"				
	/notes="vector: PTARBAC1.3; Site.1: MboI; Site.2: MboI; Hereford bull Li Domino 99375; CHORI-240 Bovine BAC library (Male) produced by Pieter de Jong"				
ORIGIN					
Query Match	8.6%;	Score 43.2;	DB 9;	Length 495;	
Best Local Similarity	50.0%;	Pred. No. 1.5;			
Matches	108;	Conservative	0;	Mismatches	108; Indels 0; Gaps 0;
QY	21	TATAACATTTTCAGTAGCATGCTGATACCAACGTTTAAACTATGGATACATATTTGAATTC	80		
DB	452	TTTAAATTTTCAAAAATGCAAAATACAAATATCTAAATTTTGAACATCTTAAT	393		
QY	81	CAAAATTTTCTTCAGATAATGTGATTTAGATTAGAGATTCAACACGGATGACACCA	140		
DB	392	CAAAAGTAACATCTGTTTAATGTGGTAATGCTGCTGGAGTCAATTTAGGTATATCA	333		
QY	141	AAGAAAACTTTGCCAAATAAGCTTTCTGTTATTTTCATAAGCAAGAGATTTAAGTTTCC	200		
DB	332	GAGAAGAATATATATAAATATGCTTTTATATGATGAGCTCTTATAGTTCCTAT	273		</

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RESULT 8
CL335225
LOCUS
DEFINITION
  CL335225
  RPCI44_253A15, r RPCI-44 Sus scrofa genomic clone RPCI44_253A15,
  genomic survey sequence.
ACCESSION
  CL335225
VERSION
  CL335225.1 GI:51387193
KEYWORDS
  GSS.
SOURCE
  Sus scrofa (pig)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
  1 (bases 1 to 618)
  Rogatcheva, M.B., Meyers, S., He, W., Larkin, D.M., Marron, B.M.,
  Beaver, J.E. and Schook, L.B.
  Piggy-BACing the Human Genome: Constructing a Porcine Physical Map
  Through Comparative Genomics
  Unpublished (2004)
  Other GSSs: RPCI44_253A15.f
  Contact: Lawrence B. Schook
  Department of Animal Sciences
  University of Illinois at Urbana Champaign
  1201 W. Gregory Dr., Urbana, IL 61801, USA
  Tel: 217 265 5326
  Fax: 217 244 5617
  Email: schook@uiuc.edu
  Clones are derived from the porcine BAC library RPCI-44
  (http://www.bacpac.chori.org/porcine242.htm). For BAC library
  availability, please contact Pieter de Jong (pdejong@chori.org).
  Clones may be purchased from BACPAC Resources
  (http://BACPACResources.chori.org). This work was undertaken as part
  of the International Swine Genome Sequencing Consortium by
  University of Illinois at Urbana Champaign, USA with funds provided
  by grant No. AG2002-34480-11928 from USDA-CSREES and
  AG2001-35205-09965 from USDA/NRI (Livestock Genome Sequencing
  Initiative)
  Plate: 253
  Seq primer: SP6
  Class: BAC ends.
  Location/Qualifiers
    1..618
    /organism="Sus scrofa"
    /mol_type="genomic DNA"
    /strain="four pigs (breed: 37.5% Yorks Landrace and 25%
    Meishan)"
    /db_xref="taxon:9823"
    /clones="RPCI44_253A15"
    /sex="male"
    /cell_type="blood"
    /note="Vector: pTARBAC2; Site 1: EcoRI; Site 2: EcoRI;
    porcine male BAC library produced by Pieter de Jong"

ORIGIN
Query Match      8.6%; Score 43.2; DB 9; Length 618;
Best Local Similarity 48.3%; Pred. No. 1.6;
Matches 117; Conservative 0; Mismatches 125; Indels 0; Gaps 0;

Qy 53 TTTAACTAGGATACATATTTGAATTCCTCAAAATTTTCTTCAGATAATGCTTAGAT 112
Db 357 TTTATTTTCAATTTTACCAATCTCTCTTCAAAAATAATACAGAGGAAATTTTATTTAAATA 416
Qy 113 TAGAGATTCAACAGGATAGACACCGAAGAAACTTTGCCAAATAAGCTTCTCGTA 172
Db 417 TGTAAATTTAGCTAAGTATAGTTTAAATTAAGTCAACAGCACTGGCCATTGAGTTA 476
Qy 173 TTTCATAGCAGAGATTTTAAGTTTTCATTTTGAAGCCATTCTGTAATTATACAACAT 232
Db 477 CTTATATAATTCAGAAAAAGCTTACTTATTTGCAATTATTACTTTTCTATCATACAGCAGA 536
Qy 233 AAAAAATGCAGTGGATATTGAACAGCTCTCTTCTGTGTAATTTCTAAATACAGTACAGTT 292

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Db 537 AAACAATGTACATATTTATCTCAAAATTTGCAAAATGCTATTCATTAANNNGTT 596
Qy 293 CA 294
Db 597 CA 598

RESULT 9
CNS0167M/c
LOCUS
DEFINITION
  CNS0167M
  Drosophila melanogaster genome survey sequence T7 end of BAC
  BACN15M24 of DrosBAC library from Drosophila melanogaster (fruit
  fly), genomic survey sequence.
ACCESSION
  AL106396
VERSION
  AL106396.1 GI:5621701
KEYWORDS
  GSS.
SOURCE
  Drosophila melanogaster (fruit fly)
ORGANISM
  Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
  Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
  Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
  1 (bases 1 to 1201)
  Genoscope.
  Direct Submission
  Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
  BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
  - Web : www.genoscope.cns.fr)
  Determination of this BAC-end sequence was carried out as part of a
  collaboration with the European Drosophila Genome Project (EDGP) -
  http://www.edgp.ebi.ac.uk -. This Drosophila melanogaster BAC
  library (Dros BAC) was made by Alain Billaud at CEPH (Centre
  d'Etude du Polymorphisme Humain) with funding provided by a MRC
  project grant. The DNA was prepared from embryos by Alain Bucheton
  and Genevieve Payan. It has been constructed in the vector
  pBeloBAC11.
  Location/Qualifiers
    1..1201
    /organism="Drosophila melanogaster"
    /mol_type="genomic DNA"
    /db_xref="taxon:7227"
    /clone="BACN15M24"
    /clone_lib="DrosBAC"
    /plasmid="pBeloBAC11"
    /note="end : T7"

ORIGIN
Query Match      8.6%; Score 43.2; DB 9; Length 1201;
Best Local Similarity 41.5%; Pred. No. 1.8;
Matches 102; Conservative 22; Mismatches 122; Indels 0; Gaps 0;

Qy 44 ATACCAAGTTTAAACATGATACATATTTGAATTCCTCAAAATTTTCTTCAGATAATGCTG 103
Db 949 AAAAAAATAATTAATAATTTTTTTTATTAATAAATAATTTTTTTTAAAAAATAAATAA 890
Qy 104 ATTAGAGATTAGAGATTCAACAGGATAGACACCGAAGAAACTTTGCCCAATAATAGC 163
Db 889 TTAATAATTTAAAAATTTTAAATATAAATAAATAAATAAATAAATAAATAAATAAATAA 830
Qy 164 TTTCTGCTATTTCATAGCAGAGATTTAAGTTTTCATTTTGAAGCCATTGTAATTA 223
Db 829 TATAAATAATTTTAAAAAATAAATAAATAAATAAATAAATAAATAAATAAATAAATAA 770
Qy 224 TACAACAATAAATAAATCAAGTGGATATTCACAGCTCTCTCTCTGTAATTTCTTAAATAC 283
Db 769 AAAAAATTAATAATAATAATTAATAATNCATAAATAAATAAATAAATAAATAAATAA 710
Qy 284 AGTACA 289
Db 709 AAAAAA 704

RESULT 10

```

```

BH742342      BH742342      623 bp      DNA      linear      GSS 25-FEB-2002
LOCUS          gu74b05.g1 BoBuds01 Brassica oleracea genomic clone gu74b05 5',
DEFINITION     genomic survey sequence.
ACCESSION      BH742342
VERSION        BH742342.1 GI:18876955
KEYWORDS       GSS.
SOURCE         Brassica oleracea
ORGANISM       Brassica oleracea
REFERENCE      Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS        Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
                rosids; eurosids I; Brassicales; Brassicaceae; Brassica.
                1 (bases 1 to 623)
                Katari,M., O'Shaughnessy,A., Palmer,L., Bahret,A., Baker,J.,
                Balija,V., Cunnius,D.M., Katzenberger,F., King,L., Kirchoff,K.,
                Kuit,K., Miller,B., Muller,S., Nascimento,L., Preston,R.,
                Santos,L., Shah,R., Zutavern,T., Dedhia,N., Rabinowicz,P.D. and
                McCombie,W.R.
                Whole Genome Shotgun Reads from Brassica oleracea (2002b)
TITLE          Unpublished (2002)
JOURNAL        Contact: W. Richard McCombie
COMMENT        Lita Annenberg Hazen Genome Sequencing Center
                Cold Spring Harbor Laboratory
                PO Box 100, Cold Spring Harbor, NY 11724, USA
                Tel: 516 367 8884
                Fax: 516 367 8874
                Email: mcombie@cshl.org
                Plate: gu74 row: b column: 05
                Seq primer: -2lUnivRev
                Class: shotgun
                High quality sequence stop: 623.
FEATURES       Location/Qualifiers
                source          1..623
                        /organism="Brassica oleracea"
                        /mol_type="genomic DNA"
                        /db_xref="taxon:3712"
                        /clones="gu74b05"
                        /clone_lib="BoBuds01"
                        /note="Vector: M13 for .x reads, pBluescript for .b and .g
                        reads; Site1: EcoRV; Whole genome shotgun library from
                        flowering buds. DNA was purified from a crude nuclear prep
                        using Brassica oleracea TO1000DH3 buds provided by Thomas
                        Osborn at the University of Wisconsin. Genomic DNA
                        provided by Pablo Rabinowicz (CSHL) and shotgun library
                        prepared in McCombie Lab."
ORIGIN
Query Match      8.5%; Score 42.6; DB 8; Length 623;
Best Local Similarity 47.5%; Pred. No. 2.3;
Matches 126; Conservative 0; Mismatches 139; Indels 0; Gaps 0;

Qy 11 ACCTTAGGATATACATTTTCAGTAGCATGCTGATACCAAGCTTTAAACTATGATACAT 70
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 192 ACCTTATAATAGCAATTTCCATAAAAATTTATTACGAAATTCGCTAATATTAGCTACCT 251
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 71 ATTGGAATCCAAATTTTCTTCAGATAATGCTGATTAGAGATTAGAGATTCAACAGGGA 130
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 252 ATAAATTTAGAGATATGCATCCTTCATTTAATTAATCAAAATCAAAATTCGATT 311
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 131 TAGACCGAAGAAAACCTTTGCCCAATAAGCTTTCTGGTATTTCATAGCAAGAGATT 190
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 312 TTGCAATCCATAAAAACATATAAAATACATTAATATTCAGATTAAATTCATTGTTTATT 371
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 191 TAAGTTTTCATTTAAGAGCCATTGTGAATTATACAAATATAAAATCAAGTGGATA 250
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 372 GAAATTAATTAATCAAAATCATTTAAATTTTACAAATATAAAATTAATCAAGTTATA 431
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 251 TTGAACAGTCTCTCTCTGATAATT 275
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 432 TTTAATTAATAGTCTCTCTTAATT 456
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 11

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```

BZ465736      BZ465736      669 bp      DNA      linear      GSS 13-DEC-2002
LOCUS          BONMQ35TF BO_1.6_2_KB_tot Brassica oleracea genomic clone BONMQ35,
DEFINITION     genomic survey sequence.
ACCESSION      BZ465736
VERSION        BZ465736.1 GI:26757807
KEYWORDS       GSS.
SOURCE         Brassica oleracea
ORGANISM       Brassica oleracea
REFERENCE      Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
AUTHORS        Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
                rosids; eurosids II; Brassicales; Brassicaceae; Brassica.
                1 (bases 1 to 669)
                Town,C.D., Van Aken,S., Utterback,T., Koo,H. and Fraser,C.M.
                Whole genome shotgun sequencing of Brassica oleracea
                Unpublished (2001)
JOURNAL        Other_GSSs: BONMQ35TR
COMMENT        Contact: Chris Town
                TIGR
                9712 Medical Center Drive, Rockville, MD 20850, USA.
                Tel: 301-838-3523
                Fax: 301-838-0208
                Email: cdtown@tigr.org
                DNA is from a doubled haploid provided by Tom Osborn.
                Seq primer: TF
                Class: sheared ends.
FEATURES       Location/Qualifiers
                source          1..669
                        /organism="Brassica oleracea"
                        /mol_type="genomic DNA"
                        /strain="TO1000DH3"
                        /db_xref="taxon:3712"
                        /clone="BONMQ35"
                        /clone_lib="BO_1.6_2_KB_tot"
                        /note="Vector: pBOS1; Site 1: BstXI; 1.6-2 kb sheared
                        total DNA inserted into pBOS1 using BstXI linkers"
ORIGIN
Query Match      8.4%; Score 42.4; DB 8; Length 669;
Best Local Similarity 50.5%; Pred. No. 2.6;
Matches 103; Conservative 0; Mismatches 101; Indels 0; Gaps 0;

Qy 6  TAAGTACCTTAGGAATATACATTTTCAGTAGCATGCTGATPACCAACGTTTAAACTATGGA 65
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 451 TAGTACCTTAAATATATATTTTCTTTAAATAATCATATAAAACAAAATTTTACACTTTATA 510
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 66 TACATATTTGAATTCCAAATTTTCTTCAGATAATGTGATTAGAGATTAGAGATTCAACC 125
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 511 TACATATTTCAAATCAAAATAAATAATTTAAATTTGATTATATATCAAAAATTTGATTCAAA 570
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 126 AGGGATAGACACCGAAAGAAACTTTGCCCAATAAGCTTTCTGTTATTTTCATAAGCAAG 185
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 571 ATATACATATGTTTAAATAATAAATTTTACTAAATAATTTTCCAAATAACCATTTATAA 630
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 186 AGATTTTAAGTTTTCATTTTAAGAA 209
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 631 AAATGGTTTCAATATATATAAGAA 654
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 12
BH661305/c
LOCUS          BOHVL59TF BO_2.3_KB Brassica oleracea genomic clone BOHVL59,
DEFINITION     genomic survey sequence.
ACCESSION      BH661305
VERSION        BH661305.1 GI:18720067
KEYWORDS       GSS.
SOURCE         Brassica oleracea
ORGANISM       Brassica oleracea
REFERENCE      Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
                Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
                rosids; eurosids II; Brassicales; Brassicaceae; Brassica.
                1 (bases 1 to 818)

```

AUTHORS Town, C.D., Van Aken, S., Utterback, T., Koo, H. and Fraser, C.M.
TITLE Whole genome shotgun sequencing of *Brassica oleracea*
JOURNAL Unpublished (2001)
COMMENT Contact: Chris Town

TIGR
 9712 Medical Center Drive, Rockville, MD 20850, USA.
 Tel: 301-838-3523
 Fax: 301-838-0208

Email: cdtown@tigr.org
 DNA is from a doubled haploid provided by Tom Osborn.
 Seq primer: TP
 Class: sheared ends.

FEATURES
 source
 Location/Qualifiers
 1..818
 /organism="Brassica oleracea"
 /mol_type="genomic DNA"
 /strain="TO1000DH3"
 /db_xref="taxon:3712"
 /clone="BOHLS9"
 /clone_lib="BO_2_3_KB"
 /note="Vector: pBAC3.6; Site 1: BstXI; 2-3 kb sheared genomic DNA inserted into pBAC3.6 using BstXI linkers"

ORIGIN

Query Match 8.4%; Score 42.4; DB 8; Length 818;
 Best Local Similarity 47.7%; Pred. No. 2.7;
 Matches 124; Conservative 0; Mismatches 136; Indels 0; Gaps 0;

QY 28 TTTCAGTACGATGCTGATACCAAGCTTTAACTATGATGATACATATTTGCAATTTCCAAATTT 87
 |||||
 Db 301 TTTCAACTAACCTAAACAAATTTAACTTTATATATACATTTTCAATTTCAAATTA 242
 |||||
 QY 88 TTCTTCAGATAATGTGATTAGAGATTAGAGATTCAACGAGGATAGACACCGAAGAAAA 147
 |||||
 Db 241 TAAATCAAGTTGATTATATACAAAATTTGATTCAAAATATATATATATATATATATAT 182
 |||||
 QY 148 CTTTGCCCAATAGCTTTCTGGTATTTTCATAGCAAGAGATTTAAGTTTCCATTTAAG 207
 |||||
 Db 181 GGAATTTCTACTAAACTATTTTTCATATACCAATTAACCAATTAACCAATTAATATATAG 122
 |||||
 QY 208 AAGCATTGTGATTTATACAAATTAACCAATTAACCAATTAACCAATTAACCAATTAACCA 267
 |||||
 Db 121 AAAATATATACAGAGCCCAATTTGAATTAACCAATTAACCAATTAACCAATTAACCAATTA 62
 |||||
 QY 268 TGATAATTTCTAAATACAGTA 287
 |||||
 Db 61 TATTAAGTTTAAAAATATA 42
 |||||

RESULT 13
 AQ544187/c
 LOCUS
 DEFINITION
 RPIC1-11-326P5, TJ RPIC1-11 Homo sapiens genomic clone RPIC1-11-326P5,
 genomic survey sequence.

ACCESSION
 AQ544187
 VERSION
 AQ544187.1 GI:4868489
 KEYWORDS
 GSS.
 SOURCE
 Homo sapiens (human)

ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 766)
 Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P., and
 Venter, J.C.

REFERENCE
 AUTHORS
 Use of BAC End Sequences from Library RPIC1-11 for Sequence-Ready
 Map Building

JOURNAL
 Unpublished (1997)
COMMENT
 Other GSSs: RPIC1-11-326P5.TV
 Contact: Shaying Zhao, William Nierman, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200

Fax: 301 838 0208
 Email: hbe@tigr.org
 Clones are derived from the human BAC library RPIC1-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genet cs (info@resgen.com). BAC end search page:
 http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
 Seq primer: SP6
 Class: BAC ends.

FEATURES
 source
 Location/Qualifiers
 1..766
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="GDB:7625164"
 /db_xref="taxon:9606"
 /clone="RPIC1-11-326P5"
 /sex="Male"
 /cell_type="Lymphocytes"
 /clone_lib="RPIC1-11"
 /note="Vector: pBAC3.6; Site 1: EcoRI; Site 2: EcoRI;
 RPIC11 Human Male BAC Library"

ORIGIN

Query Match 8.4%; Score 42.2; DB 8; Length 766;
 Best Local Similarity 49.8%; Pred. No. 3;
 Matches 107; Conservative 0; Mismatches 108; Indels 0; Gaps 0;

QY 59 CTATGGATACATATTTGCAATTTCCAAATTTTCTTTCAGATAATGTGATGAGATTAGAGA 118
 |||||
 Db 526 CTATGCTGTATATAACAAATTTATTTTGTCTATATAATATACCTAGAAATATATTA 467
 |||||
 QY 119 TTCAACGAGGATAGACACCGAAGAAATTTTCCCAAAATAGCTTTCTGGTATTTTCAT 178
 |||||
 Db 466 TTTATACAAACATTTATTCATTTGGAATATGTAATCTTTTAAATAGTCTCTGAGATTAAT 407
 |||||
 QY 179 AAGCAAGAGATTTAAGTTTCCATTTAAGAGCCATTTGTAATATATACCAATTAATAA 238
 |||||
 Db 406 AATTAGCAGATCAAGGCTATAAATTTTATAGGCTCTTGATTTATATATTGATAAATTTG 347
 |||||
 QY 239 TGCAAGTGGATATTGAACAGCTCTCTCTCGATAA 273
 |||||
 Db 346 CTCCACAAATATATAACAATTTAGTGTCAATTA 312
 |||||

RESULT 14
 CG019400

LOCUS
 ZUAAC48TV ZM_3.0_4.0 KB Zea mays genomic clone ZMMBP00005G23,
 genomic survey sequence.

ACCESSION
 CG019400
 VERSION
 CG019400.1 GI:33891565
 KEYWORDS
 GSS.
 SOURCE
 Zea mays

ORGANISM

Zea mays
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD
 clade; Panicoideae; Andropogoneae; Zea.

1 (bases 1 to 874)

Whitelaw, C.A., Quackenbush, J., Van Aken, S., Utterback, T.,
 Resnick, A., Fraser, C.M., Budiman, M.A., Bedell, J.A., Rohlfing, T.,
 Citek, R.W., Numberg, A., Robbins, D. and Lakey, N.

Consortium for Maize Genomics
 Unpublished (2002)

Other GSSs: ZUAAC48TH

Contact: Cathy Whitelaw

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA
 Tel: 301-838-5843
 Fax: 301-838-0208
 Email: whitelaw@tigr.org
 Seq primer: TP
 Class: sheared ends.

```
FEATURES
source
1. .874
/organism="Zea mays"
/mol_type="genomic DNA"
/strain="B73"
/db_xref="taxon:4577"
/clone="ZMBPa0005G23"
/clone_lib="ZM_3.0.4.0.KB"
/notes="vector: pBSK-; Site_1: HincII; 3-4 kb 'unfiltered'
genomic DNA library"

ORIGIN
Query Match 8.4%; Score 42.2; DB 9; Length 874;
Best Local Similarity 50.8%; Pred. No. 3.1;
Matches 101; Conservative 0; Mismatches 98; Indels 0; Gaps 0;
Qy 84 ATTTTCTTCAGATAATGTGATTAGAGATTAGAGATTCAACACGGGATAGACACCGAAAG 143
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 389 ATTAATATTATTTCTGTTGTAATAATAATAAGAAAAAATGCGTCTTAAATATTGGAAAA 448
Qy 144 AARACTTGGCCCAATAGCTTCTCGTATTTCATACAGAGAGATTAAAGTTTTCATT 203
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 449 AATATTTTACCCCGATGTGTACGGTTATGCAATTAATATGTTATTTTCTGCT 508
Qy 204 TAAGAAGCAATGTGCAATTATACCAATAAAAAATGCAAGTGGATATTGAACAGTCTCT 263
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 509 TTATTTTCTGAAGTAATAGTAGAAATTTAATGCAATGCGAGTATATTTGTTGAT 568
Qy 264 TCTCTGATAATCTAAATA 282
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 569 ACTTAATCGCTCAATA 587

RESULT 15
CNS000B6/c
LOCUS
DEFINITION
Drosophila melanogaster genome survey sequence TET3 end of BAC #
BAC01A24 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
ACCESSION
AL063632
VERSION
AL063632.1 GI:4938680
KEYWORDS
GSS.
SOURCE
Drosophila melanogaster (fruit fly)
ORGANISM
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
1 (bases 1 to 1101)
Genoscope.
Direct Submission
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osogawa and
Aaron Mamoser in Pieter de Jong's laboratory in the Department of
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
NY. The library is named RPCI-98 and was constructed by partial
EcoRI digestion of Drosophila DNA provided by the BDGP from the
isogenic strain y2; cn bw sp, the same strain used for the BDGP's
P1 and EST libraries. A more detailed description of the library
and how to order individual BAC clones, the entire library, or
filters for hybridization from the BACPAC Resource Center can be
found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES
source
1..1101
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
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ORIGIN
Query Match 8.4%; Score 42.2; DB 9; Length 1101;
Best Local Similarity 37.1%; Pred. No. 3.3;
Matches 108; Conservative 37; Mismatches 146; Indels 0; Gaps 0;
Qy 2 AGAGTAAGTACCTTAGGAATATTAACATTTCAGTAGCATGATACCAACGTTTAAACTA 61
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 1029 AAAAAAATAAAATTTTAATAATAAAAAATTTAAATAATAATATTTTAAATAAAWATA 970
Qy 62 TGGATACATATTTGAAATTCCTTTCAGATAATGTTAGATAGAGATTAGAGATTC 121
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 969 WATWTAAWAAAAAATAAAAAAATAAAAAAATAAAATTTTAAWATAAAWATAATA 910
Qy 122 AACCCAGGGATAGACACCGAAAGAAAACTTTTCCCAATAAAGCTTTCTGTTATTTTCA 181
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 909 ATTTAAWATWAAAAAATAAAAAATTTTAAATTTTAAATTTTAAWATAAAWATA 850
Qy 182 CAAGAGATTTAAGTTTTCGATTTTAAGAGCCATTTGGAATTATACAACAATAAAAAATGC 241
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 849 AAATWAAAAAATAAATAWAAAAAATAAAWAAAAWATAAAWATAAAWATAAAWATA 790
Qy 242 AAGTGGATATTGAACAGATCTCTCTCTGATAAATTTCTAAATACAGTACAGTT 292
Dy ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 789 AATWAAAAATTTACWAAATYWAATYAAATAAAWAAAAAATAAAWATAAAWATA 739

Search completed: November 9, 2005, 04:17:50
Job time : 2602.77 secs
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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:46:48 ; Search time 101.246 Seconds
(without alignments)
7308.644 Million cell updates/sec

Title: US-09-463-542-1_COPY_1_125

Perfect score: 125

Sequence: 1 cccctgccctgccctgcc.....gcggcgagcgggcccgccagc 125

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_16Dec04.*

1: Geneseqn1980s.*

2: Geneseqn1990s.*

3: Geneseqn2000s.*

4: Geneseqn2001as.*

5: Geneseqn2001bs.*

6: Geneseqn2002as.*

7: Geneseqn2002bs.*

8: Geneseqn2003as.*

9: Geneseqn2003bs.*

10: Geneseqn2003cs.*

11: Geneseqn2003ds.*

12: Geneseqn2004as.*

13: Geneseqn2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	125	100.0	503	2	Aax19032 Human PPA
2	118.6	94.9	12548	13	Adg36487 Human aut
3	118.6	94.9	158417	13	Adg36461 Human aut
4	81.4	65.1	201	2	Aax19063 Human PPA
5	57.8	46.2	1416	8	Abz20967 Animal te
6	56.6	45.3	420	13	Acn51217 Cotton an
7	54.4	43.5	1315	2	Aaz41312 Human nor
8	54.2	43.4	200	5	Abv61578 Human pro
9	54.2	43.4	629	13	Acn54594 Cotton an
10	53	42.4	1131	12	Adq23807 Human sof
11	52.2	41.8	168	6	Abn69641 Streptoco
12	52	41.6	3163	10	Adc87060 Human GPC
13	52	41.6	28198	10	Adg37080 Mouse pla
14	51.8	41.4	344	4	AI182007
15	51.8	41.4	437	13	Acn58918 Cotton gy
16	51.4	41.1	110000	12	Adq97050 Human can
17	51.2	41.0	349	4	AI184504 Human pol
18	51	40.8	588	13	Acn54596
19	51	40.8	600	6	Abq52497 Oligonucl
20	51	40.8	600	6	Abq52496

21	51	40.8	2188	2	Aaz77506 Human ova
22	50.8	40.6	142	7	Adr41362 Human CD-
c 23	50.8	40.6	1045	6	Abi99807 Mouse isc
c 24	50.8	40.6	1286	6	Abi99656 Mouse isc
c 25	50.6	40.5	434	4	Aai84343 Human pol
c 26	50.6	40.5	3198	2	Aax02974 Human li-
c 27	50.4	40.3	594	6	Abq43958 Oligonucl
c 28	50.4	40.3	594	6	Abq43959 Oligonucl
29	50.2	40.2	1845	3	Aaz98400 Canine be
30	50.2	40.2	1845	6	Abk40732 Dog betal
c 31	50.2	40.2	43058	6	Abi64982 Lung canc
c 32	50.2	40.2	43058	6	Abi65219 Lung canc
c 33	50.2	40.2	43058	6	Abn97455 Gene #395
c 34	50	40.0	615	8	ACA23975 Prokaryot
c 35	50	40.0	1948	6	Abq69181 Listeria
c 36	50	40.0	5082	6	Abq70953 Listeria
c 37	49.8	39.8	204	13	Adr93549 Novel S.
c 38	49.8	39.8	209	13	Adr91550 Novel S.
c 39	49.8	39.8	282	13	Adr91549 Novel S.
c 40	49.8	39.8	308	13	Adr93088 Novel S.
c 41	49.8	39.8	612	13	Adr93335 Novel S.
c 42	49.8	39.8	884	8	ACC48196
c 43	49.8	39.8	1387	10	Adc87462 Human GPC
c 44	49.8	39.8	2685	6	Abk83742 Human cDN
c 45	49.8	39.8	12733	6	Abk98631 Vector pE

ALIGNMENTS

RESULT 1

AAX19032
ID AAX19032 standard; DNA; 503 BP.

XX AC AAX19032;

XX DT 13-MAY-1999 (first entry)

XX DE Human PPAR-gamma-1 proximal promoter, exon A1 and intron A1.

XX KW Human; peroxisome proliferator activated receptor gamma; PPAR-gamma;
XX KW regulatory sequence; promoter; obesity; anorexia; lipoma; cachexia;
XX KW lipodystrophy; liposarcoma; human immunodeficiency virus; HIV;
XX KW insulin resistance; non-insulin-dependent diabetes mellitus;
XX KW polycystic ovary syndrome; gastrointestinal tract; Crohn's disease;
XX KW inflammatory bowel disease; ulcerative colitis; bowel cancer; ss.

XX OS Homo sapiens.

XX FN WO9905161-A1.

XX PD 04-FEB-1999.

XX PF 24-JUL-1998; 98WO-US015411.

XX PR 25-JUL-1997; 97US-0053692P.

XX PA (LIGA-) LIGAND PHARM INC.

XX PA (INSP) INST PASTEUR.

XX PI Briggs MR, Saladin RS, Auwerx J, Pajas L;

XX DR WPI; 1999-142844/12.

XX PT Newly isolated nucleic acid comprising a control region of a human
XX PT peroxisome proliferator activated receptor (PPAR) gamma gene - useful for
XX PT identifying modulators that are useful in treating diseases associated
XX PS with abnormal levels of human PPAR-gamma gene expression.

XX PS Claim 9; Page 78; 102pp; English.

XX CC The present invention describes an isolated, purified or enriched nucleic
XX CC acid comprising a control region of a human peroxisome proliferator


```
FT variation /*tag= a  
FT replace(697,A)  
FT /*tag= b  
XX  
XX  
XX DE20206477-U1.  
XX  
XX PD 10-OCT-2002.  
XX  
XX PF 14-MAR-2002; 2002DE-02006477.  
XX  
XX PR 14-MAR-2002; 2002DE-01012560.  
XX  
XX PA 14-MAR-2002; 2002DE-02006477.  
XX  
XX (GAGB-) GAG BIOSCIENCE GMBH.  
XX  
XX WPI; 2003-048092/05.  
XX  
XX Test kit containing bovine marker DNA, useful e.g. in animal breeding and  
XX for tracking foods, comprises specified polymorphic sequences.  
XX  
XX Claim 1; Page 18; 25pp; German.  
XX  
XX The present invention relates to a new test kit comprising marker DNA,  
XX for cattle or derived meat products, which has any of the sequences shown  
XX in AB220937-AB220970, all of which contain at least one variable position  
XX (single nucleotide polymorphism, SNP), and these SNPs are tabulated. The  
XX kit is used in animal production or breeding, and to track cattle-derived  
XX foods, e.g. for consumer protection. The specified sequences contain  
XX sufficient SNPs, present at various frequencies in the cattle population,  
XX to provide, in combination, a genetic fingerprint for individual animals.  
XX The variable positions can be queried by high-throughput methods, making  
XX possible typing of the entire cattle population in the form of a digital  
XX DNA signature. The present sequence is a bovine marker DNA for use in the  
XX test kit of the invention  
XX  
XX Sequence 1416 BP; 270 A; 253 C; 540 G; 348 T; 0 U; 5 Other;  
XX  
XX Query Match 46.2%; Score 57.8; DB 8; Length 1416;  
XX Best Local Similarity 66.4%; Pred. No. 0.7;  
XX Matches 83; Conservative 0; Mismatches 42; Indels 0; Gaps 0;  
XX  
XX Qy 1 CCCTGCCCCCTGCGCCCTGCCCCACCCACCCACCCACCCACCCACCCACCCGCGCGCGC 60  
XX Db 448 CCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCC 389  
XX  
XX Qy 61 GCGCGCCCCCGCGCGCGCGCGCTCGCGCCCGACCCCGGTTCCGCGCGCGCGCGCGC 120  
XX Db 388 CCCCCGCGCGCGCGCGCGCGCCCCCCCCCCCCCCCCCTCTCCGCCCCCCCCCCCC 329  
XX  
XX Qy 121 CCAGC 125  
XX Db 328 CCGCG 324  
XX  
XX RESULT 6  
XX ACN51217/c  
XX ID ACN51217 standard; cDNA; 420 BP.  
XX  
XX AC ACN51217;  
XX  
XX DT 02-DEC-2004 (first entry)  
XX  
XX DE Cotton androecium tissue EST Clone ID: LIB3828-011-Q1-N6-H6, SEQ:5998.  
XX  
XX KW Cotton; plant; EST; expressed sequence tag; transgenic plant; androecium;  
XX variety Nucotton33B; library LIB3828; molecular tag; molecular marker;  
XX genetic mapping; molecular mapping; seed germination; plant growth;  
XX plant quality; plant yield; plant breeding; tissue printing; ss.  
XX  
XX OS Gossypium hirsutum.  
XX  
XX PN US2004123340-A1.  
XX  
XX AC
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PD 24-JUN-2004.  
XX  
XX PF 12-DEC-2001; 2001US-00021323.  
XX  
XX PR 14-DEC-2000; 2000US-0255619P.  
XX  
XX (DEIK/) DEIKMAN J.  
XX PA (PENG/) PENG P C C.  
XX PA (FING/) FINCHER K L.  
XX PA (ZIEG/) ZIEGLER T E.  
XX  
XX Deikman J, Peng PCC, Fincher KL, Ziegler TE;  
XX WPI; 2004-479808/45.  
XX  
XX New isolated nucleic acid molecule that encodes a plant protein or its  
XX fragment, useful for isolating a variety of agronomically significant  
XX genes associated with plant growth, quality or yield, and as molecular  
XX tags to map genes.  
XX  
XX Claim 1; SEQ ID NO 5998; 34pp; English.  
XX  
XX The invention relates to 17880 cotton expressed sequence tags (ESTs;  
XX ACN45220-ACN63099). The ESTs were isolated from cDNA libraries generated  
XX from primed or non-primed seeds from variety DP50B, mature seeds from  
XX variety Coker 312 Boswell 96 Field, and androecium tissue, gynoecium  
XX tissue, developing fibres, carpel walls and septa from variety  
XX Nucotton33B. The invention also relates to substantially purified  
XX proteins or their fragments encoded by nucleic acid molecules of the  
XX invention, and to transformed plants having a nucleic acid construct  
XX comprising a nucleic acid of the invention. The cotton ESTs are useful as  
XX molecular tags to isolate genetic regions, to isolate genes, to map  
XX genes, to determine gene function and to determine whether genes are  
XX members of a particular gene family. The nucleic acid molecules may be  
XX used for isolating a variety of agronomically significant genes  
XX associated with plant growth, quality, yield, and could also serve as  
XX links in metabolic and catabolic pathways. The nucleic acid molecules are  
XX also useful for identifying genes important in initiating and maintaining  
XX seed germination or that may be used to mitigate stresses encountered  
XX during seed germination. The ESTs additionally enable the acquisition of  
XX promoters and cis-regulatory elements which will be useful to express  
XX agronomically significant genes in these tissues and/or other tissues,  
XX and also permits the acquisition of molecular markers useful in breeding  
XX schemes, genetic and molecular mapping, and in cloning of agronomically  
XX significant genes. The nucleic acid molecules are further useful for  
XX detecting the expression level or pattern of a protein or mRNA and for  
XX detecting the presence or quantity of a protein by tissue printing. The  
XX present sequence represents a specifically claimed EST isolated from a  
XX cotton variety Nucotton33B androecium tissue cDNA library (LIB3828). The  
XX sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from the US  
XX patent office at seqdata.uspto.gov/sequence.html?DocID=US20040123340  
XX  
XX Sequence 420 BP; 8 A; 55 C; 342 G; 15 T; 0 U; 0 Other;  
XX  
XX Query Match 45.3%; Score 56.6; DB 13; Length 420;  
XX Best Local Similarity 69.4%; Pred. No. 1.5;  
XX Matches 77; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
XX  
XX Qy 1 CCCCCTGCCCCCTGCCCCCTGCCCCCACCACCCACCCACCCACCCACCCAGCGCGCGCGC 60  
XX Db 145 CCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCCC 86  
XX  
XX Qy 61 GCGCGCCCCCGCGCGCGCGCGCTCGCGCCCGACCCCGGTTCCGCGCGCGCGC 111  
XX Db 85 CCGCGCCCCCCCCCGCGCGCCCCCCCCCCCCCCCCCGCGCGCGCGCGC 35  
XX  
XX RESULT 7  
XX AAZ41312  
XX ID AAZ41312 standard; cDNA; 1315 BP.  
XX  
XX AC AAZ41312;
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Result No.	Score	Query Match	Length	DB	ID	Description	
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C 2	51.4	41.1	670689	4	US-09-949-016-12505	Sequence 12505, A	
C 3	51.4	41.1	670690	4	US-09-949-016-14207	Sequence 14207, A	
4	50.6	40.5	152331	3	US-09-128-153-16	Sequence 16, Appl	
5	50.2	40.2	1845	4	US-09-614-034-188	Sequence 188, App	
6	49.8	39.8	204	4	US-09-107-433-2184	Sequence 2184, App	
7	49.8	39.8	209	4	US-09-107-433-185	Sequence 185, App	
8	49.8	39.8	282	4	US-09-107-433-184	Sequence 184, App	
C 9	49.8	39.8	308	4	US-09-107-433-1723	Sequence 1723, Ap	
C 10	49.8	39.8	612	4	US-09-107-433-1970	Sequence 1970, Ap	
C 11	49.6	39.7	320	3	US-09-165-264-11	Sequence 11, Appl	
C 12	49.4	39.5	318	3	US-09-165-264-12	Sequence 12, Appl	
C 13	49.4	39.5	319	3	US-09-165-264-8	Sequence 8, Appl	
C 14	49.4	39.5	320	3	US-09-165-264-7	Sequence 7, Appl	
C 15	49.4	39.5	320	3	US-09-165-264-13	Sequence 13, Appl	
C 16	49.4	39.5	320	3	US-09-165-264-14	Sequence 14, Appl	
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18	49	39.2	11770	4	US-09-949-016-12720	Sequence 12720, A	
19	49	39.2	11770	4	US-09-949-016-13487	Sequence 13487, A	
20	49	39.2	11770	4	US-09-949-016-13488	Sequence 13488, A	
C 21	48.8	39.0	22206	4	US-09-949-016-13901	Sequence 13901, A	
C 22	47	37.6	336	4	US-09-543-681A-4150	Sequence 4150, Ap	
C 23	47	37.6	339	4	US-09-543-681A-4147	Sequence 4147, Ap	
C 24	47	37.6	483	4	US-09-543-681A-4138	Sequence 4138, Ap	
25	47	37.6	516	4	US-09-543-681A-4140	Sequence 4140, Ap	
26	47	37.6	519	4	US-09-543-681A-4145	Sequence 4145, Ap	
27	47	37.6	608	4	US-09-543-681A-4115	Sequence 4115, Ap	


```

; Patent No. 6800744
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID
; SEQUENCES RELATING TO STREPTOCOCCUS PNEUMONIAE FOR DIAGNOSTIC
; THERAPEUTICS
;
; NUMBER OF SEQUENCES: 5206
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: <Unknown>
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: <Unknown>
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,433
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/ 085131
; FILING DATE: May 12, 1998
; APPLICATION NUMBER: 60/051553
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneka
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-011
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-8277
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 1723:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 308 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Streptococcus pneumoniae
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...308
; SEQUENCE DESCRIPTION: SEQ ID NO: 1723:
;
US-09-107-433-1723

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Matches 72; Conservative 0; Mismatches 50; Indels 0; Gaps 0;

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Qy 61 GCCCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 120
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 246 CCCCCCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 187
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Qy 121 CC 122
Db ||
Qy 186 CC 185
Db ||

RESULT 10
US-09-107-433-1970/c
; Sequence 1970, Application US/09107433
; Patent No. 6800744
; GENERAL INFORMATION:

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[illegible]

[illegible][illegible]

Qy 1 CCCCTGCCCCCTGCCCCACCCCACCCCAGCGGCGCCGC 60

Matches 77; Conservative

Matches 77; Conservative

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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 22:16:08 ; Search time 645.819 Seconds
(without alignments)
7367.447 Million cell updates/sec

Title: US-09-463-542-1_COPY_1_125

Perfect score: 125

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Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST.*

1: gb_est1.*

2: gb_est2.*

3: gb_hc.*

4: gb_est3.*

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7: gb_est6.*

8: gb_gss1.*

9: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
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C 3	63.4	50.7	982	6	CD329933	AGENCOURT
C 4	63.2	50.6	425	2	BE551555	hx97f12.x
C 5	62.4	49.9	1866	3	CR622226	full-leng
C 6	62.2	49.8	996	5	BQ708087	AGENCOURT
C 7	62	49.6	1024	6	CA975437	AGENCOURT
C 8	61.4	49.1	1674	9	CL078342	CH216-149
C 9	61.2	49.0	843	9	CNS00CS1	AL059666
C 10	61	48.8	506	9	CNS027Y	Drosophil
C 11	61	48.8	1003	9	CL475036	Tetraodon
C 12	61	48.8	1285	9	AG334095	Mus muscu
C 13	60.8	48.6	908	9	CNS006B4	AGENCOURT
C 14	60.8	48.6	1134	5	BQ651163	Drosophil
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C 20	60.2	48.2	953	9	AG391885	Mus muscu
C 21	60.2	48.2	1014	9	CL492395	SAIL_566
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C 23	60.2	48.2	1065	5	BUI48615	AGENCOURT
C 24	60.2	48.2	1142	9	CL468510	SAIL_1287

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27	60	48.0	921	9	CL465012	SAIL_1232
28	59.8	47.8	902	9	CL476702	SAIL_258
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30	59.6	47.7	899	9	CL495967	SAIL_618
C 31	59.6	47.7	925	9	CNS00CAZ	AL058951
C 32	59.6	47.7	1157	4	BM466479	Drosophil
C 33	59.6	47.7	1313	4	BM552701	AGENCOURT
C 34	59.6	47.7	1372	4	BM461243	AGENCOURT
C 35	59.6	47.7	1448	8	CC220110	CH261-92F
C 36	59.4	47.5	244	9	AG081394	Pan trogl
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C 40	59.4	47.5	1056	9	AG367644	Mus muscu
C 41	59.4	47.5	1123	9	CL470578	SAIL_144
C 42	59.4	47.5	1380	4	BM806217	AGENCOURT
C 43	59.2	47.4	468	5	BQ608819	BRY_4732
C 44	59.2	47.4	613	9	CNS01HTD	Anopheles
C 45	59.2	47.4	823	2	BF866526	963070A02

ALIGNMENTS

RESULT 1
AQ893474/c
LOCUS
DEFINITION
HS_4832_A2_C06_T7A CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=4832 Col=12 Row=E, genomic survey sequence.
ACCESSION
AQ893474
VERSION
AQ893474.1 GI:6349664
KEYWORDS
GSS.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
REFERENCE
1 (bases 1 to 821)
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS
Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D., and Hood, L.
TITLE
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
JOURNAL
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE
93180589
PUBMED
10449764
COMMENT
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones may be purchased from Research Genetics (info@resgen.com).
BAC end Web Server: http://www.htsc.washington.edu
Plate: 4832 row: E column: 12
Seq primer: T7
Class: BAC ends
High quality sequence stop: 821.
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/db_xref="taxon:9606"
/clone="Plate=4832 Col=12 Row=E"
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/clone lib="CIT Approved Human Genomic Sperm Library D"
/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in E-Coli DH10B"

ORIGIN

SOURCE ORGANISM	Homo sapiens (human)	
REFERENCE	Homo sapiens	
AUTHORS	Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
TITLE	1 (bases 1 to 1866)	
JOURNAL	Li W.B., Gruber C., Jessee J. and Polayes D.	
REMARK	Full-length cDNA libraries and normalized Unpublished	
REFERENCE	Contact : Feng Liang Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com/ Invitrogen Corporation 1600 Faraday Avenue	
AUTHORS	2 (bases 1 to 1866)	
TITLE	Genoscope.	
JOURNAL	Direct Submission	
COMMENT	Submitted (20-JUL-2004) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr - Web : www.genoscope.cns.fr)	
FEATURES	1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-strand cDNA was digested with Not I and cloned into the Not I and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized. Library was constructed by Life Technologies, a division of Invitrogen.	
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	/clone="CS0DC004Y008"	
	/tissue_type="Neuroblastoma Cot 25-normalized"	
	/plasmid="pCMVSPORT_6"	
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Qy	122	CAGC 125
Db	61	CAGC 64
RESULT 6		
BQ708087		
LOCUS	BQ708087 996 bp mRNA linear EST 16-JUL-2002	
DEFINITION	AGENCOURT.9475084 NIH_MGC_113 Homo sapiens cDNA clone IMAGE:6301404 5', mRNA sequence.	
ACCESSION	BQ708087	
VERSION	BQ708087.1 GI:21846986	
KEYWORDS	EST.	
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	1 (bases 1 to 996)	
TITLE	NIH-MGC http://mgi.nci.nih.gov/.	
JOURNAL	National Institutes of Health, Mammalian Gene Collection (MGC)	
COMMENT	Unpublished (1999)	
	Contact: Robert Strausberg, Ph.D.	
	Email: csapbs-r@mail.nih.gov	
	Tissue Procurement: Dr. Mark Watson	
	cDNA Library Preparation: Rubin Laboratory	
	DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)	
	DNA Sequencing by: Agencourt Bioscience Corporation	
	Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:	
	http://image.llnl.gov	
	Plate: LLCM2517 row: d column: 13	
	High quality sequence stop: 276.	
FEATURES	Location/Qualifiers	

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:52:53 ; Search time 4078.01 Seconds
(without alignments)
9232.381 Million cell updates/sec

Title: US-09-463-542-34_COPY_368_1144

Perfect score: 777
Sequence: 1 tcattgttagtaagactgtgt.....gtctctttttttattgttaag 777

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.*
1: gb_ba.*
2: gb_hcg.*
3: gb_in.*
4: gb_on.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	765.8	98.6	135675	9	AC093174 Homo sapi
2	765.8	98.6	148828	9	AX157024 Homo sapi
3	765.8	98.6	180963	9	AC091492 Homo sapi
C 4	765.8	98.6	221402	2	AC015995 Homo sapi
5	764.2	98.4	851	9	AF548352 Homo sapi
C 6	565.4	72.8	186028	2	AC151849 Callithri
7	421	54.2	65160	2	AC135179 Homo sapi
8	130.6	16.8	212216	2	AC131898 Oryctolag
9	103.6	13.3	185737	2	AC125447 Mus muscu
C 10	88.6	11.4	191540	2	AC136055 Rattus no
C 11	88.6	11.4	245724	2	AC091418 Rattus no
C 12	88.6	11.4	250169	2	AC120668 Rattus no
13	68.6	8.8	154195	2	AC142242 Ateles
14	53	6.8	183103	5	BX465190 Zebrafish
15	52.8	6.8	145160	2	CR847935 Danio rer
16	51.8	6.7	166021	2	AC117751 Mus muscu
C 17	50.8	6.5	153477	2	AC006278 Plasmodi
C 18	50.8	6.5	251551	3	AE014844 Plasmodi
19	50	6.4	5771	6	AX344664 Sequence

20	50	6.4	125020	9	AF429315 Homo sapi
C 21	49.6	6.4	95996	8	ATF18P9 Arabidops
22	49.4	6.4	175188	2	BX897715 Danio rer
23	49.2	6.3	18683	6	AX281291 Sequence
24	49.2	6.3	18683	6	AX345214 Sequence
C 25	49.2	6.3	156276	2	CR769765 Danio rer
C 26	49.2	6.3	175234	5	BX547997 Zebrafish
C 27	49.2	6.3	241364	2	BX571898 Danio rer
28	49	6.3	11422	6	AX345121 Sequence
29	49	6.3	11422	6	AX348323 Sequence
30	49	6.3	138121	9	AC114806 Homo sapi
31	48.8	6.3	110000	2	Continuation (9 of
32	48.8	6.3	163443	2	AC006280 Plasmodi
33	48.8	6.3	196149	2	AC004709 Plasmodi
C 34	48.8	6.3	252650	3	AE014847 Plasmodi
35	48.6	6.3	179824	9	AC096708 Homo sapi
36	48.4	6.2	13123	6	AX281322 Sequence
37	48.4	6.2	13123	6	AX344419 Sequence
38	48	6.2	92725	9	AC010350 Homo sapi
39	48	6.2	152706	9	AC008568 Homo sapi
C 40	48	6.2	152707	9	AC008876 Homo sapi
41	48	6.2	349980	6	AX344565 Sequence
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43	47.6	6.1	958	3	NAM556113 Necator a
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ALIGNMENTS

RESULT 1
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LOCUS Homo sapiens chromosome 3 clone RP11-167M22 map 3p, complete
DEFINITION AC093174 135675 bp DNA linear PRI 08-NOV-2002
AC093174
VERSION AC093174.2 GI:24796717
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 135675)
AUTHORS Wu, Q., Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J.,
Ding, H., Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D.,
Guo, Z., He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C.,
Li, P., Li, G., Li, J., Li, L., Li, S., Li, T., Liu, Y., Liu, N., Liu, B.,
Liu, Y., Li, W., Li, W., Li, X., Luo, J., Luo, Y., Qi, Q., Qi, X., Song, L.,
Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J.,
Wang, J., Wang, J., Wang, L., Wang, L., Wang, R., Wang, X., Wang, X.,
Wang, Y., Wu, D., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B.,
Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M.,
Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N.,
Yu, J. and Yang, H.

Chromosome 3p genomic sequence

Unpublished
2 (bases 1 to 135675)
AUTHORS Bao, J., Bao, Q., Bao, W., Bian, X., Cao, T., Chen, C., Chen, J., Ding, H.,
Dong, W., Fan, H., Feng, X., Gong, J., Guan, Q., Gu, X., Guo, D., Guo, Z.,
He, L., Hu, S., Huang, F., Jin, Y., Kang, N., Li, C., Li, C., Li, F.,
Li, G., Li, J., Li, L., Li, S., Li, T., Liu, Y., Liu, N., Liu, B., Liu, Y.,
Liu, W., Li, W., Li, Y., Luo, J., Luo, Y., Qi, Q., Qi, X., Song, L.,
Song, S., Sun, M., Sun, W., Sun, Y., Tan, X., Tao, R., Wang, H., Wang, J.,
Wang, J., Wang, L., Wang, L., Wang, R., Wang, X., Wang, X., Wang, Y.,
Wu, D., Wu, Q., Xie, F., Xuan, Z., Xue, Y., Yan, C., Yang, X., Yu, B.,
Zeng, Y., Zhang, G., Zhang, H., Zhang, H., Zhang, L., Zhang, M.,
Zhang, X., Zhang, X., Zhang, Y., Zhang, Y., Zhang, Z., Zhu, B., Zhu, N.,
Yu, J. and Yang, H.

Direct Submission
Submitted (13-AUG-2001) Human Genomic Center, Institute of
Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing
100101, P.R.China

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REFERENCE
AUTHORS
3 (bases 1 to 135675)
Wu,Q., Bao,J., Bao,Q., Bao,W., Bian,X., Cao,T., Chen,C., Chen,J.,
Ding,H., Dong,W., Fan,H., Feng,X., Gong,J., Guan,Q., Gu,X., Guo,D.,
Guo,Z., He,L., Hu,S., Huang,F., Jin,Y., Kang,N., Li,C., Li,C.,
Li,F., Li,G., Li,J., Li,L., Li,S., Li,T., Liu,Y., Liu,N., Liu,B.,
Lu,Y., Li,W., Li,Y., Li,Y., Luo,J., Liu,Y., Qi,Q., Qi,X., Song,L.,
Song,S., Sun,M., Sun,W., Sun,Y., Tan,X., Tao,R., Wang,H., Wang,J.,
Wang,J., Wang,L., Wang,L., Wang,R., Wang,X., Wang,X., Wang,Y.,
Wu,D., Xie,F., Xuan,Z., Xue,Y., Yan,C., Yang,X., Yu,B., Zeng,Y.,
Zhang,G., Zhang,H., Zhang,H., Zhang,L., Zhang,M., Zhang,X.,
Zhang,X., Zhang,Y., Zhang,Y., Zhang,Z., Zhu,B., Zhu,N., Yu,J. and
Yang,H.
Direct Submission
Submitted (08-NOV-2002) Human Genomic Center, Institute of
Genetics, Chinese Academy of Sciences, Datun Road, Beijing, Beijing
100101, P.R.China
On Nov 8, 2002 this sequence version replaced gi:15148930.
-----Genome Center
Center:Beijing Center
Center code:Beijing
Website:http://hgsc.igtp.ac.cn
http://www.genomics.org.cn
Contact:hgsc@igtp.ac.cn
-----Project Information
Center project name:1k project
Center clone name: RP11-167M22
-----Summary Statistics
Sequencing vector: pUC18; 100% of reads
Chemistry: Dye-terminator; Et 55% of reads
Assembly program: Big Dye; 45% of reads
Consensus quality: Phrap; version 0.990329
Consensus quality: 0 bases at least Q40
Consensus quality: 0 bases at least Q30
Consensus quality: 6 bases at least Q20
Insert size: 3392; sum-of-contigs
Quality coverage: 0.00x in Q20 bases;sum-of-contigs
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Best Local Similarity 99.1%; Pred. No. 6.5e-160;
Matches 770; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
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Qy 241 TTTGCCCTTGGCTAAGTGGTGGGACACAACTGACTGTACCTTAGGCTTAATAAACCAT 300
Db 42068 TTTGCCCTTGGCTAAGTGGTGGGACACAACTGACTGTACCTTAGGCTTAATAAACCAT 42127
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RESULT 2
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LOCUS Homo sapiens peroxisome proliferative activated receptor gamma
DEFINITION (PPARG) gene, complete cds.
ACCESSION AV157024
VERSION AV157024.2 GI:48762804
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 148828)
Rieder,M.J., da Ponte,S.H., Kuldane,K., Rajkumar,N., Smith,J.D.,
Toth,E.J. and Nickerson,D.A.
AUTHORS Direct Submission
TITLE Submitted (30-SEP-2002) Genome Sciences, University of Washington,
JOURNAL 1705 NE Pacific, Seattle, WA 98195, USA
REFERENCE 2 (bases 1 to 148828)
Rieder,M.J., Daniels,R.L., da Ponte,S.H., Hastings,N.C.,
Ahearn,M.O., Rajkumar,N., Yi,Q. and Nickerson,D.A.
AUTHORS Direct Submission
TITLE Submitted (16-JUN-2004) Genome Sciences, University of Washington,
JOURNAL 1705 NE Pacific, Seattle, WA 98195, USA
REMARK Sequence update by submitter
COMMENT On Jun 16, 2004 this sequence version replaced gi:23953882.
To cite this work please use: SeattleSNPs. NHLBI Program for
Genomic Applications, UW-FHCRC, Seattle, WA
(URL:http://pga.mbt.washington.edu).
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/organism="Homo sapiens"
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COMMENT On Nov 8, 2002 this sequence version replaced g1:13794231.
-----Genome Center
Center:Beijing Center
Center code:Beijing
Website:http://hg.igtp.ac.cn
http://www.genomics.org.cn
Contact:hg@igtp.ac.cn
----- Project Information
Center project name:RP11-33519
----- Summary Statistics
Sequencing vector: pUC18; 100% of reads
Chemistry: Dye-terminator; ET 5% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 369 bases at least Q40
Consensus quality: 571 bases at least Q30
Consensus quality: 755 bases at least Q20
Insert size: 773; sum-of-contigs
Quality coverage: 1.48x in Q20 bases; sum-of-contigs

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Best Local Similarity 99.1%; Pred. No. 6.3e-160;
Matches 770; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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Qy 301 GTGTCACTAGAGAAGTATTATTTTAAAGAGTGTGTTTGGCCATGTATAAATTTTC 360
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Qy 481 GTGATGGAATAGGAAGTAGTGAAGTGAATTTTAATAGATGTTTCTTTTATGAATAAT 540
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Qy 721 GTACACATTCGAAACATGTGTATATTTGAAAACTTCTCTCTTTTATTGTTAAG 777
Db 162568 GTACACATTCGAAACATGTGTATATTTGAAAACTTCTCTCTTTTATTGTTAAG 162624

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DEFINITION Homo sapiens chromosome 3 clone RP11-33519 map 3, WORKING DRAFT
SEQUENCE, 36 unordered pieces.
AC015995
VERSION AC015995.5 GI:9966961
KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 221402)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.
TITLE Homo sapiens chromosome 3, Clone RP11-33519
JOURNAL Unpublished
2 (bases 1 to 221402)

REFERENCE 1 (bases 1 to 221402)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, M., Baldwin, J., Barna, N., Becker, R., Boguslavskiy, L., Boukhgalter, B., Brown, A., Castelle, A., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dearellano, K., Dewar, K., Domino, M., Donelan, L., Doyle, M., Ferreira, P., Fitzhugh, W., Forrest, C., Funke, R., Gage, D., Galagan, J., Gardyna, S., Grant, G., Hagos, B., Heaford, A., Horton, L., Howland, J., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J., Lehoczy, J., Liu, C., Locke, K., Macdonald, P., Marquis, N., McSwan, P., McGurk, A., McKernan, K., McLaughlin, J., Meldrum, J., Morrow, J., Naylor, J., Norman, C. H., O'Connor, T., O'Donnell, P., Peterson, K., Pollara, V., Riley, R., Roy, A., Santos, R., Severy, P., Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J., Tesfaye, S., Tirrell, A., Vassiliev, H., Vo, A., Wheeler, J., Wu, X., Wyman, D., Ye, W. J., Zimmer, A. and Zody, M.
Direct Submission
Submitted (18-NOV-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 221402)

REFERENCE 1 (bases 1 to 221402)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S., Barna, N., Bastien, V., Boguslavskiy, L., Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Karatas, A., LaRocque, K., Lamazares, R., Landers, T., Lehoczy, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Marquis, N., Matthews, C., McCarthy, M., McSwan, P., McKernan, K., McPheters, R., Meldrum, J., Meneus, L., Mihova, T., Mieng, V., Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Riback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R., Seaman, S., Severy, P., Sougne, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Travets, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (24-AUG-2002) Whitehead Institute/MIT Center for Genome

TITLE
JOURNAL


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Best Local Similarity 99.1%; Pred. No. 6.1e-160;
Matches 770; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Qy 1 TCATGTAGCTAAGACTGTGTAGAAATGTCGGGTCTCGATGTTGGCGCTATTCAAGCCCTGA 60
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Db 97993 ACTTCTAGAGATAGAAAGCTTGTAGGCTCAGGAAAGCAAACTTCAAGATGAAATCC 97934
Qy 661 AATAGAGAGCTTAATTTTATTTGGCATGTACATTTTGGCAGCTTGTGCTGCTGCTGCA 720
Db 97933 AATAGAGAGCTTAATTTTATTTGGCATGTACATTTTGGCAGCTTGTGCTGCTGCTGCA 97874
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Db 97873 GTACACATCTGAACTGTGTATTTATTTGAAATCTTCTCTCTCTCTCTCTCTCTCTCTGCA 97817

RESULT 5
AF548352
LOCUS
DEFINITION Homo sapiens peroxisome proliferator-activated receptor gamma 3
AF548352
ACCESSION
VERSION AF548352.1 GI:28626258
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 851)
AUTHORS Meirhaeghe,A., Fajas,L., Helbecque,N., Vertebrate; Euteleostomi;
Auwerx,J. and Amouyel,P. Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.
TITLE A functional polymorphism in a STAT5B site of the human PPAR gamma
3 gene promoter affects height and lipid metabolism in a French
```

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population
JOURNAL Arterioscler. Thromb. Vasc. Biol. (2003) In press
REFERENCE 2 (bases 1 to 851)
AUTHORS Meirhaeghe,A., Fajas,L., Helbecque,N., Auwerx,J. and Amouyel,P.
TITLE Direct Submission
JOURNAL Submitted (23-SEP-2002) INSERM U508, Institut Pasteur de Lille, 1
rue du Pr Calmette - BP 245, Lille 59019, France
FEATURES
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/db_xref="taxon:9606"
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3"
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ORIGIN
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Best Local Similarity 99.0%; Pred. No. 3.1e-159;
Matches 769; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

Qy 1 TCATGTAGGTAAAGACTGTGTAGATGTCGGGTCTCGATGTTGGCGCTATTCAAGCCCTGA 60
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Qy 61 TGATAAGGCTTTTGGCATTAGATGCTGTTTGTCTTCATGAAATACAGACTATTCCTAGG 120
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Qy 121 ATCTTGGAGCTTTTCATAGAGATAAGCTTGTGAATCTTAAGAGCTTGTGCTGCA 180
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Qy 361 AAACATTAACTTTTCAGGCTTATATTTTAAAGGATCGTTTTCGCTGCTGCTGCTGCA 420
Db 361 AAACATTAACTTTTCAGGCTTATATTTTAAAGGATCGTTTTCGCTGCTGCTGCTGCA 420
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481 GTGATGAAATAGGAAAGTAGGTGAAGTGAATTTAATAGATGTTCTTTTATGAATAAT 540
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RESULT 6

AC151849/c
LOCUS AC151849 186028 bp DNA linear HTG 08-OCT-2004
DEFINITION Callithrix jacchus clone CH259-42F14, WORKING DRAFT SEQUENCE, 2
ordered pieces.

AC151849

AC151849.1 GI:53983836

HTG; HTGS_PHASE2; HTGS_DRAFT.

KEYWORDS Callithrix jacchus (white-tufted-ear marmoset)

SOURCE Callithrix jacchus

ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Platyrrhini; Callitrichidae;
Callithrix.

REFERENCE 1 (bases 1 to 186028)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 186028)

AUTHORS Cheng, J.-F., Hamilton, M., Peng, Y., Mukherjee, S., Hosseini, R.,

Peng, Z., Malinov, I. and Rubin, E.M.

TITLE Direct Submission

JOURNAL Submitted (08-OCT-2004) Genome Sciences, Lawrence Berkeley National
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA

COMMENT

Sequence Produced by Berkeley PGA

Web site: <http://pga.lbl.gov>

Center Code: PGABERK

Center Project Name: J108-42F14

Bac Clone Name: CH259-42F14

This sequence has been compared to sequences of other species
using VISTA (<http://www-gsd.lbl.gov/VISTA>). The results can be
viewed at:
http://pga.lbl.gov/cgi-bin/search_cvcgd?type=ncvalue=PPARG

The order-orientation of the draft sequence was accomplished by
using:
Avid (<http://baboon.math.berkeley.edu/avid>) and paired end
information.

Funding agent: Programs for Genomic Applications (NHLBI)

Summary Statistics:

Sequencing vector: Plasmid; pUC18

Chemistry: Dye-terminator Big Dye

Assembly program: Phrap version 0.990329.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 2 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces

* is believed to be correct as given, however the sizes

* of the gaps between them are based on estimates that have

* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 26679: contig of 26679 bp in length
* 26680 26779: gap of unknown length
* 26780 186028: contig of 159249 bp in length.

FEATURES

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/organism="Callithrix jacchus"
/mol_type="genomic DNA"
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/clone="CH259-42F14"

ORIGIN

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Best Local Similarity 87.8%; Pred. No. 1.8e-115;
Matches 663; Conservative 0; Mismatches 86; Indels 6; Gaps 4;
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Qy 138 AAGAGATAAGGTTGTGNAATCCTAAGACCCCTAGGACCAATTACTTAGATGATCTGCTCTCT 197
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123485 AAGAGATAAGGTTGTGAATCCTAAGACCCCTGGGCCCTTTTACTTAGGCAATCTGTTCTCT 123426
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123425 TGCTCATC---TGAAAGTCTGCTTCGTAAGTGTACACATGCAATTTGCCCTTGCCTACATG 123369
Qy 258 G-TGTGSCACACAACTGTACTGTCACCTTAGGCTTAATAACCAATGTCATCTAGAAATGA 316
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123368 GTTGAGGCACACATCTGTACTGTCTATTTTAGGCTTAATAACCAATGTCATCTAGAAATGA 123309
Qy 317 AGTTATATTTTAAAGGATCGTTTGGCCATGATATAAATTTTCAACATTAACCTTCAG 376
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123308 AGTTATATTTTAAAGGATTTGTTTGGCCGTGATTAAGTTATCACAATTAACCTTCGA 123249
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Qy 497 AGTAGGTGAAGTGAATTTTAATAGATGTTCTTTTATGAATAAATTTTAAAGATTTGCC 556
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122949 GTATTATGATATACATA-TTTGGCAGCTAGGTTGTATATGATGTACAAATCTGAGCA 122891
Qy 737 TGTGTGTATATTGAAAATCTTGTCTCTTTTATT 771
Db |||||
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RESULT 7

AC135179
LOCUS Homo sapiens chromosome 17 clone RP11-1200B1 map 17, LOW-PASS linear HTG 10-OCT-2002
SEQUENCE SAMPLING.
AC135179
VERSION AC135179.2 GI:23683246
KEYWORDS HTG; HTGS_PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
1 (bases 1 to 65160)
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Birren, B., Nusbaum, C. and Lander, E.
JOURNAL Unpublished
TITLE Homo sapiens chromosome 17, clone RP11-1200B1
AUTHORS 2 (bases 1 to 65160)
Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B.,
Canarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.,
Cook, A., Cooke, P., DeArelano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,
Karatas, A., Kelle, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Miho, T., Miengo, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Tefaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.

TITLE Direct Submission
JOURNAL Submitted (08-OCT-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
AUTHORS 3 (bases 1 to 65160)
Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B.,
Canarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.,
Cook, A., Cooke, P., DeArelano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,
Karatas, A., Kelle, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Miho, T., Miengo, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schuback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Tefaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.

TITLE Direct Submission
JOURNAL Submitted (10-OCT-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT On Oct 10, 2002 this sequence version replaced gi:23592138.
All repeats were identified using RepeatMasker:
Smit, A.P.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIER
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L28344
Center clone name: 1200_B_1

* NOTE: This record contains 80 individual

* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.

1 720: contig of 720 bp in length
2 820: gap of 100 bp
3 821 1524: contig of 704 bp in length
4 1525 1624: gap of 100 bp
5 1625 2325: contig of 701 bp in length
6 2326 2425: gap of 100 bp
7 2426 3211: contig of 786 bp in length
8 3212 3311: gap of 100 bp
9 3312 4038: contig of 727 bp in length
10 4039 4138: gap of 100 bp
11 4139 4869: contig of 731 bp in length
12 4870 4969: gap of 100 bp
13 4970 5704: contig of 735 bp in length
14 5705 5804: gap of 100 bp
15 5805 6538: contig of 734 bp in length
16 6539 6638: gap of 100 bp
17 6639 7365: contig of 727 bp in length
18 7366 7465: gap of 100 bp
19 7466 8184: contig of 719 bp in length
20 8185 8284: gap of 100 bp
21 8285 8988: contig of 704 bp in length
22 8989 9088: gap of 100 bp
23 9089 9806: contig of 718 bp in length
24 9807 9906: gap of 100 bp
25 9907 10615: contig of 709 bp in length
26 10616 10715: gap of 100 bp
27 10716 11447: contig of 732 bp in length
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32 13074 13173: gap of 100 bp
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34 13892 13991: gap of 100 bp
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62 25498 26104: contig of 607 bp in length

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* 26915	27014: gap of 100 bp	bp in length
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* 27747	27846: gap of 100 bp	bp in length
* 27847	28566: contig of 720 bp	bp in length
* 28567	28666: gap of 100 bp	bp in length
* 28667	29399: contig of 733 bp	bp in length
* 29400	29499: gap of 100 bp	bp in length
* 29500	30225: contig of 726 bp	bp in length
* 30226	30325: gap of 100 bp	bp in length
* 30326	31052: contig of 727 bp	bp in length
* 31053	31152: gap of 100 bp	bp in length
* 31153	31871: contig of 719 bp	bp in length
* 31872	31971: gap of 100 bp	bp in length
* 31972	32686: contig of 715 bp	bp in length
* 32687	32786: gap of 100 bp	bp in length
* 32787	33508: contig of 722 bp	bp in length
* 33509	33608: gap of 100 bp	bp in length
* 33609	34328: contig of 720 bp	bp in length
* 34329	34428: gap of 100 bp	bp in length
* 34429	35148: contig of 720 bp	bp in length
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* 39333	40060: contig of 728 bp	bp in length
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* 42563	43291: contig of 729 bp	bp in length
* 43292	43391: gap of 100 bp	bp in length
* 43392	44102: contig of 711 bp	bp in length
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* 44203	44934: contig of 732 bp	bp in length
* 44935	45034: gap of 100 bp	bp in length
* 45035	45722: contig of 688 bp	bp in length
* 45723	45822: gap of 100 bp	bp in length
* 45823	46553: contig of 731 bp	bp in length
* 46554	46653: gap of 100 bp	bp in length
* 46654	47365: contig of 712 bp	bp in length
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* 48195	48294: gap of 100 bp	bp in length
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Matches 430; Conservative 0; Mismatches 15; Indels 0; Gaps 0;		
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Qy 393	AAGTCTAGTTTTTCTTAAGTCTGCGATGATAGAGTATCGTCATTCATGACATAA	452
Db 33693	AAGTCTAGTTTTTCTTAAGTCTGCGATGATAGAGTATCGTCATTCATGACATAA	33752
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Db 33873	TGATGAATCAATTTTGCTGCTGTAGTTACTTTTAGAGAATAGAAGCATTTAGGCTCA	33932
Qy 633	GGAAAGCAACATTCAGAATCAATCAATAGAGAAGGTAAATTTATTTGGGCATGTAC	692
Db 33933	TGAAAGCAACATTCAGAATCAATCAATAGAGAAGGTAAATTTATTTGGGCATGTAC	33992
Qy 693	ATTTTGGCAGCTAGGCTGTACATGTGTACATTTCTGAACATGTGTGTATATTGAAA	752
Db 33993	ATTTTGGCAGCTAGGCTGTACATGTGTACATTTCTGAACATGTGTGTATATTGAAA	34052
Qy 753	ATCTTGTCTCTTTTATTGTTAAG 777	
Db 34053	ATCTTGTCTCTTTTATTGTTAAG 34077	
RESULT 8		
AC131898		
LOCUS		
DEFINITION		
Oryctolagus cuniculus clone LBI-83M7, WORKING DRAFT SEQUENCE, 12		
unordered pieces.		
AC131898		
AC131898.1 GI:22507060		
VERSION		
HTG; HTGS_PHASE1; HTGS_DRAFT.		
KEYWORDS		
Oryctolagus cuniculus (rabbit)		
SOURCE		
Oryctolagus cuniculus		
ORGANISM		
Mammalia; Euthera; Chordata; Craniata; Vertebrata; Euteleostomi;		
Eukaryota; Metazoa; Lagomorpha; Leporidae; Oryctolagus.		
REFERENCE		
1 (bases 1 to 212216)		
AUTHORS		
Martin, J., Schwartz, J.R., Hosseini, R., Peng, Y., Peng, Z., Rubin, E.M.		
and Cheng, J.-F.		
TITLE		
Direct Submission		
JOURNAL		
Unpublished		
REFERENCE		
2 (bases 1 to 212216)		
AUTHORS		
Martin, J., Schwartz, J.R., Hosseini, R., Peng, Y., Peng, Z., Rubin, E.M.		
and Cheng, J.-F.		
TITLE		
Submitted (27-AUG-2002) Genome Sciences, Lawrence Berkeley National		
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA		
JOURNAL		
Draft Sequence Produced by Berkeley PGA		
COMMENT		
Web site: http://pga.lbl.gov		
Center Code: PGABERK		
Center Project Name: R052-83M7		
Bac Clone Name: LBI-83M7		
Additional information on comparative analysis and ordering are		
available at:		
http://pga.lbl.gov/cgi-bin/search/cvcd?type=n&value=PPARG		
Funding agent: Programs for Genomic Applications (NHLBI)		
Contact: 'Jody Schwartz' jrschwartz@lbl.gov		
if library name is LBI to LB4, please see website		
for a description: http://www-gsd.lbl.gov/cheng/BAC.html		
Summary Statistics:		
Sequencing vector: Plasmid; pUC18		
Chemistry: Dye-terminator Big Dye		
Assembly: This is a 'working draft' sequence. It currently		
consists of 12 contigs. The true order of the pieces		
is not known and their order in this sequence record is		
arbitrary. Gaps between the contigs are represented as		
runs of N, but the exact sizes of the gaps are unknown.		
This record will be updated with the finished sequence		
as soon as it is available and the accession number will		
be preserved.		
1		
2430: contig of 2430 bp in length		
2431		
2530: gap of unknown length		
2531		
4596: contig of 2066 bp in length		
4597		
4696: gap of unknown length		
4697		
9216: contig of 4520 bp in length		

```
* 9217 9316: gap of unknown length
* 9317 15539: contig of 6223 bp in length
* 15540 15539: gap of unknown length
* 15640 21088: contig of 5449 bp in length
* 21089 21188: gap of unknown length
* 21189 33983: contig of 12795 bp in length
* 33984 34083: gap of unknown length
* 34084 48193: contig of 14010 bp in length
* 48194 48193: gap of unknown length
* 48194 67641: contig of 19448 bp in length
* 67642 67741: gap of unknown length
* 67742 91391: contig of 23650 bp in length
* 91392 91491: gap of unknown length
* 91492 117177: contig of 25686 bp in length
* 117178 117277: gap of unknown length
* 117278 144157: contig of 26880 bp in length
* 144158 144257: gap of unknown length
* 144258 212216: contig of 67959 bp in length.
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    /mol_type="genomic DNA"
    /db_xref="taxon:9986"
    /clone="LBI-83W7"
ORIGIN
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Best Local Similarity 62.5%; Pred. No. 4.8e-19;
Matches 307; Conservative 0; Mismatches 144; Indels 40; Gaps 5;
QY 287 GCCTTAATACCATGTCATCTAGATGAAGTATATATTTAAAGAGTCGTTTTCGCC 346
Db 84838 GGCCTTAACCAACCCATCGCGCTGATGCAATTTATA-TTCAGGAACATATATCTATGAC 84896
QY 347 ATGTATATAATTTTCAACATTAACCTTCAGCGTATTATATCTTTTAAAGGCTAGTTTTT 406
Db 84897 TCTGTTGATCATC-----TGCCTTTCCAGATATTAATCTTTTAAAGATCTAGTTTTT 84950
QY 407 CTTAAGTCTGTGAGTATAGAGTATCGTCAATTCATGTGACATAAAGATGGAAGGGG 466
Db 84951 CTTAAGTCTGCATATATATATATATGATGATGATTAATTTGGCATAAAGATGGATGAT 85010
QY 467 CTTCAATTCATGTTAGTATGGAATAGGAATAGGATGATGATTTTAAATAGATGTTTC 526
Db 85011 AAAGACCTTGCACTCACAATGATGATGATGATGATGATGATGATGATGATGATGAT 85070
QY 527 TTTTATGAATAATTTTAAAGATGTCAGCGCTGATGATGATGATGATGATGATGATGAT 586
Db 85071 CTCATAAAATCAATTTAAGAGATGATTTGCAATCCCTGATGATGATGATGATGATGAT 85130
QY 587 GTGGTCTGTTAGTACTTTTATAGAGATAGAAAGCATTTAGGCTCAGGGAAGCAAAACAT 646
Db 85131 GTGGCTCTGTTAGTACTTTATAGAGCAAAAG-----CAT 85166
QY 647 TCAGATGAATCCAAATGAGAGGTAATTTATTTGGGCAATGATACATTTTGGCGAGCCTA 706
Db 85167 TCAGATGAATCCAAACAAAGGA-AAAAATTTATTTGATGATGATGATGATGATGATGAT 85225
QY 707 GGCTGTGTACATGTTACACATCTCGACATGTTGATATTTGAAATCTGTCCTCTTTT 766
Db 85226 G-----ATTGCTAAGGATTTTAAATATGTTGATGATGATGATGATGATGATGATGAT 85277
QY 767 TTAATGTTAAG 777
Db 85278 GAAATGTTTCAG 85288
```

```
RESULT 9
AC125447
LOCUS
DEFINITION Mus musculus chromosome 6 clone RP24-507D15 map 6, *** SEQUENCING
IN PROGRESS ***, 3 unordered pieces.
ACCESSION AC125447
```

VERSION
KEYWORDS
SOURCE
ORGANISM

AC125447.4 GI:45237223
HTG; HTGS PHASE1; HTGS FULLTOP; HTGS_ACTIVEFIN.
Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Bixren,B., Nusbaum,C. and Lander,E.
Mus musculus chromosome 6, clone RP24-507D15

REFERENCE

AUTHORS

TITLE

JOURNAL

AUTHORS

REFERENCE

AUTHORS

2 (bases 1 to 185737)
Bixren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hages,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kellis,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrum,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Riese,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tefaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.

TITLE

JOURNAL

REFERENCE

AUTHORS

Direct Submission
Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 185737)

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

Bixren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,
Anderson,M., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,
Boguslavskiy,L., Boukhalter,B., Camarata,J., Chang,J., Choepel,Y.,
Collymore,A., Cook,A., Cooke,P., Corum,B., DeArelano,K.,
Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S.,
Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S.,
Graham,L., Grand-Pierre,N., Hages,B., Hagen,N., Hagopian,D., Hages,B.,
Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kellis,C., Landers,T., Levine,R.,
Lindblad-Toh,K., Liu,X., Liu,A., Mabbitt,R., Maclean,C.,
Macdonald,P., Major,J., Manning,J., Mlenga,T., Mihova,T., Murphy,T., Naylor,J.,
Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,
O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Riese,C., Rogov,P.,
Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C.,
Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,
Talamas,J., Tefaye,S., Theodore,J., Topham,K., Travers,M.,
Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X.,
Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

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REFERENCE

AUTHORS

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REFERENCE

AUTHORS

Submitted (06-MAR-2004) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 6, 2004 this sequence version replaced gi:29029334.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L26714
Center clone name: 507_D_15

* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as

- * runs of N, but the exact sizes of the gaps are unknown.
- * This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

* 1 54649: contig of 54649 bp in length
* 54650 54749: gap of 100 bp
* 54750 156594: contig of 101845 bp in length
* 156595 156694: gap of 100 bp
* 156695 185737: contig of 29043 bp in length.

FEATURES

Source

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/mol_type="genomic DNA"
/db_xref="taxon:10090"
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/map="6"
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ORIGIN

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Matches 225;	Conservative 0;	Mismatches 159;	Indels 6;	Gaps 2;
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Db	129614	TTTGGAGTTGCTAATCCCTTTTAAANAATCTAGTCTTTCATAGTCTGAATACTGATATAAT	129673	
QY	431	TATCGTCA--TTTCATGTGCATAAAGAATGAAAGGGCTTCATTCATGTTAGTGTATGGA	488	
Db	129674	TATTAATTAACCTGATACTAAAAATGGGTATAATAGAGTTTTCATCATGTCAGTGACCCCT	129733	
QY	489	AATAGGAAAGTAGGTGAAGTGAATTTTAATAGATGT-----TTCCTTTATGAAATAATTTTT	544	
Db	129734	ACAGGTAAACAAGAGAAGTGGATCAACAGCTATCTGCTTCTTTTATGCAACTACTCTG	129793	
QY	545	AAAAGATTTGTCAGCGCCCTGCATGATTATATGATGAATCATTTTGTGGTCTCTTAGTTACTTT	604	
Db	129794	ATGAGAGTTTGGGTACTGTGTGATCCATGATGAGTCGTTTAATGGTGATATCGTTATTT	129853	
QY	605	TTAGAGAAATAGAAAGCATTTGTAGGCTCAGGGAAGCAAAACATTTCAGAAATGAAATCCAAATA	664	
Db	129854	CTAGAGAAATATAAAGTTTTATATGCTCAGGGAAGCAAAATACTCGGGATGAAAAATTAATG	129913	
QY	665	GAGAAGGTAAATTTTATTTTGGGCATGTACATTTTGGCAGCCTAGCGTGTGACATGTGTAC	724	
Db	129914	TGTAATTTTGCATTTTGGCAGATTGGAAATGTCAGAAATTTAGAAATCAACAAATTTTTC	129973	
QY	725	ACATTCTCGAACATGTGTGTATATTGAAAT	754	
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RESULT	10
AC136055/c	
LOCUS	
DEFINITION	AC136055 191540 bp DNA linear HTG 29-OCT-2002 Rattus norvegicus clone CH230-123L10, *** SEQUENCING IN PROGRESS ***, 62 unordered pieces.
ACCESSION	AC136055
VERSION	AC136055.1 GI:24417926
KEYWORDS	HTG; HTGS PHASE1.
SOURCE	Rattus norvegicus (Norway rat)
ORGANISM	Rattus norvegicus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE

1. (bases 1 to 191540)

Muzny, D., Marie, Metzger, M. Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Baldwin, D., Bandaranaike, A., Ayodeji, M., Baca, E., Baden, H., Salwaybech, V., Koyagi, A., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhat, C., Burch, P., Burrell, K., Calderon, E.,

Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., B. Eaves, K., Egan, A., Escoto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Frazer, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Geiger-Georgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guveara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hoques, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowals, C., Kratt, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorusluhewa, L., Loulseged, H., Lozada, R. J., Lu, X., Ma, J., Maheshwari, M., Mahindaratne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhney, S., Mcleod, M., McNeill, T., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Mundisasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokemelehen, O., Okuonwu, G., Olarunlunsaqoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Puzos, M., Quito, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S., J. Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Smajds, D., Sneed, A., Sodergruen, E., Song, X., Z., Sorelle, R., Soza, J., Steinkle, M., Strong, R., Sutton, A., Svatek, A., Taber, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villaseña, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wlezyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weis, R., Smith, D. R., Holt, R. A., Smith, H. O., Weinstein, G., and Gibbs, R. A.

TITLE
JOURNAL.

JOURNAL
REFERENCE
AUTHORS
Unpublished
2 (bases 1 to 191540)
Rat Genome Sequencing Consortium.

TITLE Direct Submission
JOURNAL Submitted (29-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

COMMENT
----- Genome Center
Center: Baylor College of Medicine

Web site: http://www.hasc.bcm.tmc.edu/cancer_code_bcm

web site: <http://www.hgsa.com>
 Contact: hgac=hel m@hgsa.com

----- Project Information -----
Contact: ngyuc-het@comp.cmc.edu

----- Project Information
Context project name: KDDC

Center project name: KUDG
Center clone name: CH230-123110

Center clone name: CH230-I23L

----- Summary Statistics -----

Sequencing vector: Plasmid;

Chemistry: Dye-terminator Big Dye: 100% o

Assembly program: Phrap; version 0.990329

Consensus quality: 138097 bases at least Q40

Consensus quality: 143025 bases at least Q30

Consensus

★ NOTE: Estimated insert size may differ from sequence length
 ★ (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 62 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N , but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1135: contig of 1135 bp in length
* 1236: gap of unknown length
* 1236: contig of 1449 bp in length
* 2685: gap of unknown length
* 2785: contig of 1303 bp in length
* 4088: gap of unknown length
* 4188: contig of 1414 bp in length
* 5602: gap of unknown length
* 5702: contig of 1483 bp in length
* 7185: gap of unknown length
* 7285: contig of 1305 bp in length
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* 10204: gap of unknown length
* 10304: contig of 1604 bp in length
* 11908: gap of unknown length
* 12008: contig of 1025 bp in length
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* 14693: gap of unknown length
* 14793: contig of 1205 bp in length
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* 28390: gap of unknown length
* 28490: contig of 1323 bp in length
* 29913: gap of unknown length
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* 31237: gap of unknown length
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* 34429: contig of 2583 bp in length
* 37012: gap of unknown length
* 37112: contig of 2649 bp in length
* 39761: gap of unknown length
* 39861: contig of 1457 bp in length
* 41318: gap of unknown length
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* 44118: contig of 1604 bp in length
* 45721: gap of unknown length
* 45821: contig of 1387 bp in length
* 47208: gap of unknown length
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* 49261: contig of 3289 bp in length
* 52551: gap of unknown length
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* 56585: gap of unknown length
* 58264: contig of 1680 bp in length
* 58365: gap of unknown length
* 61441: contig of 3076 bp in length
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* 61541: contig of 2076 bp in length
* 63617: gap of unknown length
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* 70926: gap of unknown length
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* 74192: contig of 2263 bp in length
* 76455: gap of unknown length
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* 79102: contig of 3380 bp in length
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* 98678: gap of unknown length
* 98778: contig of 5301 bp in length
* 104079: gap of unknown length
* 104179: contig of 3868 bp in length
* 108047: gap of unknown length
* 108147: contig of 4239 bp in length
* 112386: gap of unknown length
* 112486: contig of 3698 bp in length
* 116184: gap of unknown length
* 116284: contig of 6471 bp in length
* 122755: gap of unknown length
* 122855: contig of 191540; DB 2; Length 191540;
Query Match 11.4%; Score 88.6; DB 2; Length 191540;
Best Local Similarity 60.7%; Pred. No. 1e-09;
Matches 184; Conservative 0; Mismatches 109; Indels 10; Gaps 2;

Qy 371 TTTTCAGGGTTATTAACTCTTTTAAAGTCTAGTCTTTTCTTAAAGTCTGTCAGTAAATAGG 430
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Db 13920 TATTATTAAATTGATATTAAATAATGGACATAATAAAGACATTTGTCATCATGTCAGTGA 13861
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Db 13860 CCCTAGTGGTAAACAAAGAGAGAGTGAATCAATACCTATCTGTCTCTTTTATGCAACCAT 13801
Qy 541 TTTTAAAGATTGTCAGCCCTGTCATGATTTATGATGAATCATTTTGGTCTGTAGTT 600
Db 13800 TCTGAGGAGAGTTGGGTGTCAGATGATCCAAACATGATGATCATTTAGTGTGTGTGTG 13741
Qy 601 ACTTTTAGAGAAATAGAAAGCATTTAGGCTCAGGGAAGCAACATTCAGATGAAATCC 660
Db 13740 ATTTCTAGAGATATAAAGCTTTACATGCTCGGAGAAAGCAATCTGTGGATGAAAT 13681
Qy 661 AAT 663
Db 13680 AAT 13678

RESULT 11
AC091418/c

LOCUS AC091418 245724 bp DNA linear HTG 13-NOV-2002
DEFINITION Rattus norvegicus clone CH230-2G6, *** SEQUENCING IN PROGRESS ***.
ACCESSION AC091418
VERSION AC091418.5 GI:24941354
KEYWORDS HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_ENRICHED.

SOURCE
ORGANISM
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
AUTHORS
1 (bases 1 to 245724)
Muzny, D.M., Adams, C., Adio-Oduola, B., Ali-osman, F.R., Allen, C.,
Albrooks, S.L., Amarantunga, H.C., Are, J.R., Ayale, M., Banks, T.,
Barbaria, J., Benton, J., Bmaga, K., Blankenburg, K., Bonnin, D.,
Bouck, J., Bowie, S., Brieva, M., Brown, E., Brown, M., Bryant, N.P.,
Buhay, C., Burch, P., Burrell, C., Burrell, K.L., Byrd, N.C.,
Carroll, T.P., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D.,
Chen, G., Chen, R., Chen, Z., Chowdhry, I., Christopoulos, C.,
Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R.,
Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A.,
Delaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H.,
Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J.,
Earhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Escotto, M.,
Falls, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P., Frantz, P.,
Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N., Gill, R.,
Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S., Hamilton, K.,
Harris, C., Harris, K., Hart, M., Haviak, P., Hawes, A., Hernandez, J.,
Hernandez, O., Hodgson, A., Hogues, M., Holloway, C., Hollins, B.,
Honsi, F., Howard, S., Huber, J., Huly, S., Hume, J., Jackson, L.E.,
Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S.,
Karleson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C.,
Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lewis, L.C., Lewis, L.,
Li, J., Li, Z., Licharge, O., Lieu, C., Liu, C., Liu, J., Liu, W., Louissegh, H.,
Lozado, R.J., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J.,
Maheshwari, M., Mapua, E., Martin, R., Martindale, A., Martinez, E.,
Massey, E., Mawhney, E., McLeod, M.P., Meador, M., Mei, G., Metzker, M.,
Miner, G., Miner, Z., Mitchell, T., Mohabbat, K., Morgan, M., Morris, S.,
Moser, M., Neal, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N.,
Nguyen, N., Nickerson, E., Nwokwenkwo, S., Ogih, M., Okwundu, G.,
Oragunye, N., Oviedo, R., Pace, A., Payton, B., Peery, J., Perez, L.,
Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y.,
Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savery, G.,
Scherer, S., Scott, G., Shen, H., Shooshitari, N., Sison, I.,
Sodergren, E., Sonake, T., Sparks, A., Stanley, H., Stone, H.,
Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H.,
Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S.,
Umaney, K., Vasquez, L., Vera, V., Villalon, D., Vinson, R., Wang, Q.,
Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S.,
Williams, G., Williamson, A., Wleczyk, R., Wooden, S., Worley, K.,
Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,
Weinstock, G. and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 245724)
Worley, K.C.

Direct Submission
Submitted (20-APR-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 245724)
Worley, K.C.

Direct Submission
Submitted (13-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA

COMMENT
On Nov 13, 2002 this sequence version replaced gi:22855703.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: TUOY
Center clone name: CH230-2G6
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 220652 bases at least Q40
Consensus quality: 223895 bases at least Q30
Consensus quality: 225929 bases at least Q20
Estimated insert size: 229375; sum-of-contigs estimation
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 245724: contig of 245724 bp in length.

FEATURES
source
1. .245724
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-266"
misc_feature
1. .1001
/notes="wgs_contig"
misc_feature
243338..245724
/notes="wgs_contig"

ORIGIN
Query Match 11.4%; Score 88.6; DB 2; Length 245724;
Best Local Similarity 60.7%; Pred. No. 9.8e-10;
Matches 184; Conservative 0; Mismatches 109; Indels 10; Gaps 2;
Qy 371 TTTCAGGGTTATTAACTCTTTTAAAGGCTAGTTCTTCTTAAGTCTGTGCGAGTAATAGAGG 430
Db 52722 TTTCAGGCTTCTAATCCCTTTAAAGATCTAGGTTTCTTAAGTCTGAATGGTAATGTAAT 52663
Qy 431 TATCGTCATTCATGTGCATATAAAG-----ATGGAAGGGGCTTCATTTCATGTTAGTGA 484
Db 52662 TATTATTATTTGATATTAAAAAATGGACATAATAAAGACTTTTGCATCATGTCAGTGA 52603
Qy 485 TGGAAATAGAGAGTAGGTGAAGTGAATTTAATAGA-----TGTTCTTTTATGAAATAAT 540
Db 52602 CCCTAGTGGTAAAAACAGAGAGAGTGAATCAATACCTATCTCTCTCTTTTATGCAACCAT 52543
Qy 541 TTTTAAAGATGTCGCCAGCCCTGCATGATTATTATGATGAATCATTTCTGGTCTCTTAGTT 600
Db 52542 TCTGAGAGAGAGTTTGGGTGCTGCATGATCCAAAGATGATCAATTTAGTGGTGTGTTGGTG 52483
Qy 601 ACTTTTAGAGATAGAAAGCATTTGTAGGCTCAGGGAAGCAAAATTCAGAAATCAATATCC 660
Db 52482 ATTCTAGAGATATAAAGCTTTTACATGCTCGGAGAAAGCAAAATCTCTGATGATAAATT 52423
Qy 661 AAT 663
Db 52422 AAT 52420

RESULT 12
AC120668/c AC120668 250169 bp DNA linear HTG 23-NOV-2002
LOCUS Rattus norvegicus clone CH230-24K10, *** SEQUENCING IN PROGRESS
DEFINITION

[illegible]

```
Best Local Similarity 60.7%; Pred. No. 9.8e-10;
Matches 184; Conservative 0; Mismatches 109; Indels 10; Gaps 2;

Qy 371 TTTGAGGTTATTAACTCTTTTAAGTCTAGTCTTTCTTAAGTCTGCGAGTATAGAGG 430
Db 191916 TTTGAGCTGCTAAATCCCTTTTAAAGATCTAGGTTTCTTAAGTCTGAATGTAATGTAAT 191857
Qy 431 TATCGTCATTCATGTGACATATAAG-ATGAAAGGGCTTCATTCATGTTAGTGA 484
Db 191856 TATTATTAAATTTGATATTAAAAATGGACATAATAAAGACTTTTGCATCATGTCAGTGA 191797
Qy 485 TGGAAATAGGAAGTAGTGAAGTGAATTTTAATAGA---TGTTTCTTTTATAGAAATTAAT 540
Db 191796 CCCTAGTGGTAAAAACAAGAGAAGTGAATCAATACCTATCTGCTCTCTTTTATGCAACCAT 191737
Qy 541 TTTTAAAGATGTCAGCCCTGTCATGATTTATCATGAATCATTTTCTGCTGCTGTTAGTT 600
Db 191736 TCTGAGGAGAGTTTGGGTGCTGATGATCCAACTGATGATCATTTAGTGGTGTGTTGGTG 191677
Qy 601 ACTTTTAGAGTAAGAAGCAATTTAGCTCAGGAAAGCAACATTCAGAAATGAAATCC 660
Db 191676 ATTTCTAGAGATATAAAGCTTTACATGCTCGGAGAAGCAATACTGTGATGAATTT 191617
Qy 661 AAT 663
Db 191616 AAT 191614

RESULT 13
AC142242 154195 bp DNA linear HTG 07-AUG-2003
LOCUS Atelerix albiventris clone LB4-81H3, WORKING DRAFT SEQUENCE, 5
DEFINITION ordered pieces.
ACCESSION AC142242
VERSION AC142242.2 GI:33469169
KEYWORDS HTG; HTGS PHASE2; HTGS DRAFT.
SOURCE Atelerix albiventris (Middle-African hedgehog)
ORGANISM Atelerix albiventris
Mammalia; Eutheria; Insectivora; Erinaceidae; Erinaceinae;
Atelerix.
REFERENCE 1 (bases 1 to 154195)
AUTHORS Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
Direct Submission
Unpublished
2 (bases 1 to 154195)
AUTHORS Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
Direct Submission
Submitted (27-MAR-2003) Genome Sciences, Lawrence Berkeley National
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
3 (bases 1 to 154195)
AUTHORS Cheng,J.-F., Hamilton,M., Peng,Y., Mukherjee,S., Hosseini,R.,
Peng,Z., Malinov,I. and Rubin,E.M.
Direct Submission
Submitted (07-AUG-2003) Genome Sciences, Lawrence Berkeley National
Laboratory, 1 Cyclotron Rd., Berkeley, CA 94720, USA
On Aug 7, 2003 this sequence version replaced gi:29294044.
```

Sequence Produced by Berkeley PGA
Web site: <http://pga.lbl.gov>
Center Code: PGABERK
Center Project Name: E015
Bac Clone Name: LB4-81H3

This sequence has been compared to sequences of other species using VISTA (<http://www-gsd.lbl.gov/VISTA>). The results can be viewed at:
http://pga.lbl.gov/cgi-bin/search_cvcgd?type=n&value=PPARG

The order-orientation of the draft sequence was accomplished by using:

Avid (<http://baboon.math.berkeley.edu/mavid>),
Lagan (<http://lagan.stanford.edu/>) and paired end information.

Funding agent: Programs for Genomic Applications (NHLBI)

If the Bac Library Name is LB1 to LB4, please see website for the description: <http://www-gsd.lbl.gov/cheng/BAC.html>
These libraries are available through the BACPAC Resources Center: <http://www.chori.org/bacpac/libraryres.htm> as LBNU-1 to LBNU-4.

Summary Statistics:
Sequencing vector: Plasmid; pUC18
Chemistry: Dye-terminator Big Dye
Assembly program: Phrap version 0.990329.
* NOTE: This is a 'working draft' sequence. It currently consists of 5 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and the accession number will be preserved.
* 1 33275: contig of 33275 bp in length
* 33276 33375: gap of unknown length
* 33376 54251: contig of 20876 bp in length
* 54252 54351: gap of unknown length
* 54352 65733: contig of 11382 bp in length
* 65734 65833: gap of unknown length
* 65834 127515: contig of 61682 bp in length
* 127516 127615: gap of unknown length
* 127616 154195: contig of 26580 bp in length.

FEATURES
source
1..154195
/organism="Atelerix albiventris"
/mol_type="genomic DNA"
/db_xref="taxon:9368"
/clone="LB4-81H3"

ORIGIN

Query Match 8.8%; Score 68.6; DB 2; Length 154195;
Best Local Similarity 62.3%; Pred. No. 2.9e-05;
Matches 124; Conservative 0; Mismatches 74; Indels 1; Gaps 1;
Qy 558 GCCTGTCATGATTTATGATGAATCATTTTGTGGTCTCTTAGTCTACTTTTACAGAAATAGAA 617
Db 17762 GTCTGTAAATTCACACGAGGTCACCTGCTAGTCTGTTAGTCTCTCTGCGGAACAA 17821
Qy 618 AGCATTTAGGCTCAGGAAAGCAACATTCAGAAATCCAAATAGAGAGGTAAATT 677
Db 17822 AGCATGATAAACACAGGGGACAGAG-CACCTCAGTGTGAAGTACAGCAGCAAAAACAAGT 17880
Qy 678 TATTGGGCATGTACATTTTGGCAGCCTAGCTGTGTACATGTGTACACATTCGACAT 737
Db 17881 TAGTTGTGCTGAACATTTTGGCAGCCTTGGTGTCTTAAGACTCTGAATGTTAAAAATG 17940
Qy 738 GTGTGTATATTGAAAAATCT 756
Db 17941 TTGTTCTTTTACITTT 17959

RESULT 14
BX465190 183103 bp DNA linear VRT 13-NOV-2003
LOCUS BX465190
DEFINITION Zebrafish DNA sequence from clone DKEY-121A11 in linkage group 22,
complete sequence.
ACCESSION BX465190
VERSION BX465190.4 GI:38323074
KEYWORDS HTG.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Cypriniformes; Cyprinidae; Danio.
1 (bases 1 to 183103)
Barlow, K.
Direct Submission
Submitted (13-NOV-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfsh-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Nov 13, 2003 this sequence version replaced gi:32567584.
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: <http://www.sanger.ac.uk>
Contact: zfsh-help@sanger.ac.uk

During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest, except on the rare
occasion of the clone being a YAC.

The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
EM, EMBL; SW, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information

http://www.sanger.ac.uk/Projects/C_elegans/wormpep Clone-derived
Zebrafish pUC subclones occasionally display inconsistency over the
length of mononucleotide A/T runs and conserved TA repeats. Where
this is found the longest good quality representation will be
submitted.

Repeat names beginning 'Dr' were identified by the Recon repeat
discovery system (Zhifeng Bao and Sean Eddy, submitted), and those
beginning 'drr' were identified by Rick Waterman (Stephen Johnson
lab, WashU). For further information see
http://www.sanger.ac.uk/Projects/D_rerio/fishmask.shtml DXEY-121A11
is from a Zebrafish BAC library

VECTOR: pIndigoBAC-5.

FEATURES
source
1..183103
Location/Qualifiers
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clones="DXEY-121A11"
/clone_lib="DanioKey"

ORIGIN
Query Match 6.8%; Score 53; DB 5; Length 183103;
Best Local Similarity 48.5%; Pred. No. 0.081;
Matches 176; Conservative 0; Mismatches 185; Indels 2; Gaps 1;
QY 321 ATATTTTAAAGGATCGTTTTCGCCATGTATATAATTTTCAACATTAACATTTTCAGGGTT 380
DB 159530 ATATGTTTAACTACTATTACCTGACGTGTATACATTTTTCMAATATTTTACACACATTTG 159589
QY 381 ATTAATCCTTTTAAGGCTAGTGTCTTTTCTTAAGTCTGTGCGAGTAATAGAGGT--ATCGTCA 438
DB 159590 TTTTAAACCTAAAAGTTCTCTATACTATTGTGTTTTTGGCCCTGATTACAGTACATCATAT 159649
QY 439 TTTATGTGACATAAAGATGGAAGGGCTTCATTCATGTAGTGATGGAATAGGAAG 498
DB 159650 TTTACTAGTATTTTAGAAGATACTAGTATTTCAGCTTAAAGTGAATTTGAATGGTTTAAAT 159709
QY 499 TAGGTGAAGTCATTTTAAATAGATGTTCTTTTATGAATAATTTTAAAGATGTCCAG 558
DB 159710 TAGGTGAAGTCATTTGACACACACTGGTCTGTTGTGCAACCATCGAATAAATAATTTGAT 159769

QY 559 CCTGCGATGATTTATGATGAATCATTTTGTGGTCTGTGTAGTACTTTTAGAGATAGAAA 618
DB 159770 AATATTGACCATATTGACCTGGAAAATTAATTTAAAAAATTTAATTACAGTCAGCTCTAAA 159829
QY 619 GCATTGTAGCTCAGGGAGAGCAACATTCAGATGAATCCCAATAGAGAGGTAAATTTT 678
DB 159830 AGAATAAGACTGAATAGAAAATAGACAGAGAAGAAAACATTAAATACCGTAAATTC 159889
QY 679 ATT 681
DB 159890 CTT 159892

RESULT 15

CR847935

LOCUS

DEFINITION

CR847935

VERSION

CR847935.2

KEYWORDS

HTG; HTGS PHASE1

SOURCE

ORGANISM

Danio rerio

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Cypriniformes; Cyprinidae; Danio.

1 (bases 1 to 145160)

Sims, S.

Direct Submission

Submitted (10-OCT-2004) Wellcome Trust Sanger Institute, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquiries:

zfsh-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk

On Oct 11, 2004 this sequence version replaced gi:54019745.

----- Genome Center

Center: Wellcome Trust Sanger Institute

Center code: SC

Web site: <http://www.sanger.ac.uk>

Contact: zfsh-help@sanger.ac.uk

----- Project Information

Center project name: zC194N10

----- Summary Statistics

Assembly program: XGAP4; version 4.5

Chemistry: Dye-terminator; 100% of reads

Consensus quality: 141804 bases at least Q40

Consensus quality: 142048 bases at least Q30

Consensus quality: 142375 bases at least Q20

Insert size: 144560; sum-of-contigs

Insert size: 150827; 3.5% error; agarose-fp

Quality coverage: 11.77x in Q20 bases; sum-of-contigs Quality

coverage: 11.28x in Q20 bases; agarose-fp

* NOTE: This is a 'working draft' sequence. It currently

* consists of 7 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 47977: contig of 47977 bp in length

* 47978 48077: gap of 100 bp

* 48078 61854: contig of 13777 bp in length

* 61855 61954: gap of 100 bp

* 61955 114748: contig of 52794 bp in length

* 114749 114749: gap of 100 bp

* 114849 119330: contig of 4482 bp in length

* 119331 119430: gap of 100 bp

* 119431 127887: contig of 8457 bp in length

* 127888 127888: gap of 100 bp

* 127988 134017: contig of 6030 bp in length

* 134018 134117: gap of 100 bp

* 134118 145160: contig of 11043 bp in length.

* Location/Qualifiers

FEATURES

```
source
1. .145160
/organism="Danio rerio"
/mol_type="genomic DNA"
/db_xref="taxon:7955"
/clone="CH211-194N10"
/clone_lib="CHORI-211"
1. .47977
/note="assembly fragment:00776"
fragment_chain:1
clone_end:T7
vector_side:left"
48078. .61854
/note="assembly fragment:00558"
fragment_chain:1"
61955. .114748
/note="assembly fragment:01633"
fragment_chain:2"
114849. .119330
/note="assembly fragment:00004"
fragment_chain:2"
119431. .127887
/note="assembly fragment:00176"
fragment_chain:2"
127988. .134017
/note="assembly fragment:00071"
fragment_chain:2"
134118. .145160
/note="assembly fragment:00347"
fragment_chain:2
clone_end:SP6
vector_side:right"
```

ORIGIN

```
Query Match          6.8%; Score 52.8; DB 2; Length 145160;
Best Local Similarity 44.9%; Pred. No. 0.093;
Matches 201; Conservative 0; Mismatches 247; Indels 0; Gaps 0;

Qy 320 TATATTTTAAAGGATCGTTTTTGGCCATGATATAAATTTTCAAAACATTAACCTTCAGGGT 379
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 9856 TTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTATTTAT 9915
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 380 TATTAATCTTTTAAAGCTAGTTTTTCTTAAAGCTCTGCGAGTAATAGAGGTATCGTCAT 439
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 9916 TATTATTATTATTATTATTATTATTATTATTATTATTATTATTATTATTATTATTATTAG 9975
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 440 TCATGTGACATAAAGATGGAAGGGCTTCATTTCATGTTAGTCATGGAATAGGAAAGT 499
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 9976 TATTATTATTAGTATTATTATCATTTATTTATTATTATTATTATTATTATTACACTTTC 10035
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 500 AGGTGAAGTGATTTTAAATAGATGTTCTTTTATGAAATAATTTTAAAGAGATTGCCAGC 559
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 10036 ACTGTGCTGCTGTTATTTAAATATTTATTTATTTATTTATTTATTTATTTATTTATTT 10095
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 560 CCTGCATGATTTATGATGAATCATTTTGTGCTGCTGTTAGTTACTTTTGTAGAAATAGAAAG 619
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 10096 ATTATTATTATTATTATAAATGTTTATTGCTTGTGTTACGAACGTTTAAAGTGAAAGTC 10155
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 620 CATTGTAGGCTCAGGAAAGCAACATTCAGAATGAAATCCAATGAGAGGTAATTTA 679
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 10156 TTTCAATTATGCTGTCATATTAGCTATTTTCCCTTATATAAATAAACAATAAATAA 10215
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 680 TTTGGGCATGTACATTTTGGCAGCCTAGGCTGTGTACATGTGTACACATTTCTGAACATGT 739
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 10216 TAAAACTTTAAATATGTTAAGAAATATGACCCCTTCTCTGCACAGCAAGTCAAAAATAAA 10275
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 740 GTGTATATTGAAATCTTGTCTCTTTT 767
Db      |||||      |||||      |||||      |||||      |||||      |||||
Qy 10276 TTGTATATTGTCAAAATGTCTTTTTT 10303
Db      |||||      |||||      |||||      |||||      |||||      |||||
```

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:46:48 ; Search time 629.342 Seconds
(without alignments)
7308.644 Million cell updates/sec

Title: US-09-463-542-34_COPY_368_1144

Perfect score: 777

Sequence: 1 tcattgtagtaagactgtgt.....gtctctttttattgttaag 777

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_16Dec04.*
1: Geneseq1980s.*
2: Geneseq1990s.*
3: Geneseq2000s.*
4: Geneseq2001as.*
5: Geneseq2001bs.*
6: Geneseq2002as.*
7: Geneseq2002bs.*
8: Geneseq2003as.*
9: Geneseq2003bs.*
10: Geneseq2003cs.*
11: Geneseq2003ds.*
12: Geneseq2004as.*
13: Geneseq2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	777	100.0	1463	2	AAX19034
2	764.2	98.4	158417	13	ADS36461
3	383	49.3	468	2	AAX19065
4	200.6	25.8	201	13	ADS39355
5	200.6	25.8	201	13	ADS39165
6	196.6	25.3	201	13	ADS39169
7	195.8	25.2	201	13	ADS39163
8	50	6.4	5771	6	ABN80072
9	49.2	6.3	18683	6	ABL32312
10	49.2	6.3	18683	6	ABL54333
11	49	6.3	11422	6	ABK39937
12	49	6.3	11422	6	ABL32219
13	48.4	6.2	13123	6	ABK31423
14	48.4	6.2	13123	6	ABL54364
15	47	6.0	6880	6	ABK31322
16	47	6.0	6880	6	ABL70293
17	47	6.0	6880	6	ABL70293
18	46.8	6.0	6179	4	AAS46343
19	46.8	6.0	6179	4	ABK31250
20	46.6	6.0	33053	6	ABQ67006

C	21	46.4	6.0	50000	6	ABL56202	ABL56202	AmEPV gen
	22	45.2	5.8	2000	6	ABZ17494	ABZ17494	Arabidops
	23	45.2	5.8	5718	4	AAS46464	AAS46464	Tumour su
	24	45.2	5.8	5718	6	ABL33373	ABL33373	Human imm
	25	44.4	5.7	5886	6	ABL34213	ABL34213	Human imm
	26	44.4	5.7	5992	6	AAS61208	AAS61208	Human gen
	27	44.4	5.7	16811	6	ABL33946	ABL33946	Human imm
	28	44.4	5.7	18624	6	ABL33702	ABL33702	Human imm
	29	44.2	5.7	5937	6	ABL34542	ABL34542	Human met
	30	44.2	5.7	5937	6	ABN80150	ABN80150	Human che
	31	44.2	5.7	5937	7	ADS99803	ADS99803	Bisulphit
	32	44.2	5.7	6057	6	ABK31397	ABK31397	Signal tr
	33	44.2	5.7	6057	6	ABL70362	ABL70362	Chemical
	34	44.2	5.7	6823	6	ABL33145	ABL33145	Human imm
	35	44	5.7	13584	6	ABL32615	ABL32615	Human imm
	36	43.8	5.6	17918	6	AAS61418	AAS61418	Human gen
	37	43.6	5.6	69727	10	ACF65374	ACF65374	Phototrab
	38	43.6	5.6	110000	10	ACF67367	ACF67367	Continuation (36 o
	39	43.4	5.6	2000	6	ABZ16087	ABZ16087	Arabidops
	40	43.4	5.6	6849	6	ABL92253	ABL92253	Chemical
	41	43.4	5.6	10543	6	ABK31249	ABK31249	Signal tr
	42	43.4	5.6	10543	6	ABL70206	ABL70206	Chemical
	43	43.4	5.6	10543	6	AAS61161	AAS61161	Human gen
	44	43.4	5.6	34548	6	ABL70603	ABL70603	Chemical
	45	43.2	5.6	6056	6	ABL33026	ABL33026	Human imm

ALIGNMENTS

RESULT 1

AAX19034

ID AAX19034 standard; DNA; 1463 BP.

AC AAX19034;

XX

DT 13-MAY-1999 (first entry)

XX

DE Human PPAR-gamma-3 proximal promoter, exon A2 and intron A2.

XX

KW Human; peroxisome proliferator activated receptor gamma; PPAR-gamma;
KW regulatory sequence; promoter; obesity; anorexia; lipoma; cachexia;
KW lipodystrophy; liposarcoma; human immunodeficiency virus; HIV;
KW insulin resistance; non-insulin-dependent diabetes mellitus;
KW polycystic ovary syndrome; gastrointestinal tract; Crohn's disease;
KW inflammatory bowel disease; ulcerative colitis; bowel cancer; ss.

XX Homo sapiens.

XX

XX WO9905161-A1.

XX

XX 04-FEB-1999.

XX

XX 24-JUL-1998; 98WO-US015411.

XX

PR 25-JUL-1997; 97US-0053692P.

XX

PA (LIGA-) LIGAND PHARM INC.

PA (INSP) INST PASTEUR.

XX

PI Briggs MR, Saladin RS, Auwerx J, Fajas L;

XX

XX WPI; 1999-142844/12.

XX

PT Newly isolated nucleic acid comprising a control region of a human

PT peroxisome proliferator activated receptor (PPAR) gamma gene - useful for
PT identifying modulators that are useful in treating diseases associated
PT with abnormal levels of human PPAR-gamma gene expression.

XX

PS Claim 11; Page 88; 102pp; English.

XX

CC The present invention describes an isolated, purified or enriched nucleic
CC acid comprising a control region of a human peroxisome proliferator

activated receptor gamma (PPAR-gamma) gene. The nucleic acids are useful for screening for agents capable of modulating the expression of a human PPAR-gamma gene. These agents (modulators) form pharmaceutical compositions that are useful for treating diseases associated with high/low levels of human PPAR-gamma gene expression. The diseases include obesity, anorexia, cachexia, lipodystrophy, lipomas, liposarcomas, abnormalities associated with anti-human immunodeficiency virus (HIV) treatment, insulin resistance, non-insulin-dependent diabetes mellitus (NIDDM), polycystic ovary syndrome, diseases of the gastrointestinal (GI) tract, inflammatory bowel disease, Crohn's disease, ulcerative colitis and bowel cancer. The nucleic acids are useful for studying the role of the PPAR-gamma gene in various diseases and disorders. The structure of PPAR-gamma enables genetic studies of PPAR-gamma mutations in humans, and evaluation of its role in disorders like insulin resistance, NIDDM, and diseases associated with altered adipose tissue function, like obesity and lipodystrophic syndromes. The nucleic acids are also useful for gene therapy and the production of transgenic animals, which are useful in screening assays. The control regions of the nucleic acids enable screening for modulators of the human PPAR-gamma gene, which are useful in designing drugs for treating disorders or diseases associated with the level of PPAR-gamma gene expression. The present sequence represents human PPAR-gamma-3 proximal promoter, exon A2 and intron A2

Seq Sequence 1463 BP; 461 A; 228 C; 284 G; 490 T; 0 U; 0 Other;

Query Match 100.0%; Score 777; DB 2; Length 1463;
Best Local Similarity 100.0%; Pred. No. 2.4e-183;
Matches 777; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCATGTAGGTAAGACTGTGTAGAGATGCGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 60
Db 368 TCATGTAGGTAAGACTGTGTAGAGATGCGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 427

Qy 61 TGATAAGGCTTTTGGCATTAGATGCTGTTTGTCTTCATGGAATACAGCTATCTAGG 120
Db 428 TGATAAGGCTTTTGGCATTAGATGCTGTTTGTCTTCATGGAATACAGCTATCTAGG 487

Qy 121 ATCTCTGAGGCTTTTCAAGAGATAAGGTTGTGAATCTTAAGACCTAGGACCAATTTTACT 180
Db 488 ATCTCTGAGGCTTTTCAAGAGATAAGGTTGTGAATCTTAAGACCTAGGACCAATTTACT 547

Qy 181 TAGATGATCTGCTCTCTGTTGCTCTGTAAGAGTCTGTTCTGAGGGGTGCTGCA 240
Db 548 TAGATGATCTGCTCTCTGTTGCTCTGTAAGAGTCTGTTCTGAGGGGTGCTGCA 607

Qy 241 TTTGCTTGCCTAGTCTGTTGCTGCACTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
Db 608 TTTGCTTGCCTAGTCTGTTGCTGCACTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 667

Qy 301 GTGTCATCTAGATGAAGTTATATTTTAAAGAGATCGTTTTCGCCATGTATATAATTTTC 360
Db 668 GTGTCATCTAGATGAAGTTATATTTTAAAGAGATCGTTTTCGCCATGTATATAATTTTC 727

Qy 361 AAACATTAACTTTCAGGGTTAATCTTTTAAAGTCTAGTTTCTTAAAGTCTGCTGCA 420
Db 728 AAACATTAACTTTCAGGGTTAATCTTTTAAAGTCTAGTTTCTTAAAGTCTGCTGCA 787

Qy 421 GTAATAGAGGATCTGCTATCTGATGACATAAAGATGGAAGGGGCTTCAATCATCTTA 480
Db 788 GTAATAGAGGATCTGCTATCTGATGACATAAAGATGGAAGGGGCTTCAATCATCTTA 847

Qy 481 GTGATGGAATAGGAAGTAGGTGAAGTGAATTTTAAATAGATGTTTCTTTTATGAATAAT 540
Db 848 GTGATGGAATAGGAAGTAGGTGAAGTGAATTTTAAATAGATGTTTCTTTTATGAATAAT 907

Qy 541 TTTTAAAGATGTGCTCCAGCCCTGATGATTTATGATGAATCATTTTGTGCTGCTAGTT 600
Db 908 TTTTAAAGATGTGCTCCAGCCCTGATGATTTATGATGAATCATTTTGTGCTGCTAGTT 967

Qy 601 ACTTTTGAAGATAGGAAGATCTGAGCTCAGGGAAGCAACATTTTCAGATCAATCC 660
Db 968 ACTTTTGAAGATAGGAAGATCTGAGCTCAGGGAAGCAACATTTTCAGATCAATCC 1027

Qy 661 AATAGAGAGGTAAATTTATTTGGCATGTACATTTTGGCAGCCTAGGCTGTGTACATGT 720
Db 1028 AATAGAGAGGTAAATTTATTTGGCATGTACATTTTGGCAGCCTAGGCTGTGTACATGT 1087

Qy 721 GTACACATCTCGAACATGTGTGTATATATGAAATCTTTGTCTCTTTTATTTGTTAAG 777
Db 1088 GTACACATCTCGAACATGTGTGTATATGAAATCTTTGTCTCTTTTATTTGTTAAG 1144

RESULT 2
ADS36461
ID ADS36461 standard; DNA; 158417 BP.
XX AC ADS36461;
XX DT 16-DEC-2004 (first entry)
XX Human autoimmune disease-related genomic DNA sequence - SEQ ID 1675.
XX single nucleotide polymorphism detection; SNP detection;
XX rheumatoid arthritis; type 1 diabetes; multiple sclerosis;
XX systemic lupus erythematosus; inflammatory bowel disease; psoriasis;
XX thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;
XX glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;
XX primary systemic vasculitis; ds.
XX Homo sapiens.
XX OS
XX PN WO2004083403-A2.
XX PD 30-SEP-2004.
XX PF 18-MAR-2004; 2004WO-US008461.
XX PR 18-MAR-2003; 2003US-0455444P.
XX PR 25-APR-2003; 2003US-0465241P.
XX (APPL-) APPLERA CORP.
XX Cargill M, Begovich AB, Alexander HC;
XX WPI; 2004-728480/71.
XX DR
XX PT New isolated nucleic acid molecule comprises at least 8 contiguous nucleotides where one of the nucleotides is a single nucleotide polymorphism (SNP), useful for diagnosing or treating autoimmune diseases, e.g. rheumatoid arthritis.
XX PS Claim 16; SEQ ID NO 1675; 123pp; English.
XX CC The invention comprises amino acid and coding sequences containing genetic polymorphisms associated with an altered risk of developing an autoimmune disease (e.g. rheumatoid arthritis). The invention further comprises a method of identifying an individual that has an altered risk of developing an autoimmune disease, comprising detecting a single nucleotide polymorphism (SNP) in a nucleic acid of the invention. The DNA and protein sequences of the invention are useful for diagnosing and treating autoimmune diseases, such as: rheumatoid arthritis, type 1 diabetes, multiple sclerosis, systemic lupus erythematosus, inflammatory bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease, CC myocardiitis, Sjogren's disease, or primary systemic vasculitis. The present nucleic acid represents a human autoimmune disease-related genomic DNA sequence of the invention. NOTE: The present sequence is not shown in the specification, but has been retrieved from the WIPO website.
XX SQ Sequence 158417 BP; 47887 A; 30186 C; 31475 G; 48577 T; 0 U; 292 Other;
Query Match 98.4%; Score 764.2; DB 13; Length 158417;
Best Local Similarity 98.6%; Pred. No. 1.3e-179;
Matches 766; Conservative 4; Mismatches 7; Indels 0; Gaps 0;

Qy 1 TCATGTAGGTAAGACTGTGTAGAAATGTCGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 60


```
Db 300 GGCATGTACATTTTGGCAGCCTAGGTGTGTACATGTGTACACATTTCTGAACATGTGTG 359
Qy 743 TATATTGAAATCTTGTCTCTTTTATTGTTAAG 777
Db 360 TATATTGAAATCTTGTCTCTTTTATTGTTAAG 394

RESULT 4
ID ADS39355 standard; DNA; 201 BP.
XX ADS39355;
XX
DT 16-DEC-2004 (first entry)
XX
DE Human autoimmune disease-related SNP context sequence - SEQ ID 4569.
XX
KW single nucleotide polymorphism detection; SNP detection;
KW rheumatoid arthritis; type 1 diabetes; multiple sclerosis;
KW systemic lupus erythematosus; inflammatory bowel disease; psoriasis;
KW thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;
KW glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;
KW primary systemic vasculitis; ds.
XX
OS Homo sapiens.
XX
PN WO2004083403-A2.
XX
PD 30-SEP-2004.
XX
PF 18-MAR-2004; 2004WO-US008461.
XX
PR 18-MAR-2003; 2003US-0455444P.
PR 25-APR-2003; 2003US-0465241P.
XX
PA (APPL-) APPLERA CORP.
XX
PI Cargill M, Begovich AB, Alexander HC;
XX
WPI; 2004-728480/71.
XX
PT New isolated nucleic acid molecule comprises at least 8 contiguous
PT nucleotides where one of the nucleotides is a single nucleotide
PT polymorphism (SNP), useful for diagnosing or treating autoimmune
PT diseases, e.g. rheumatoid arthritis.
XX
PS Claim 16; SEQ ID NO 4569; 123pp; English.
XX
CC The invention comprises amino acid and coding sequences containing
CC genetic polymorphisms associated with an altered risk of developing an
CC autoimmune disease (e.g. rheumatoid arthritis). The invention further
CC comprises a method of identifying an individual that has an altered risk
CC of developing an autoimmune disease, comprising detecting a single
CC nucleotide polymorphism (SNP) in a nucleic acid of the invention. The DNA
CC and protein sequences of the invention are useful for diagnosing and
CC treating autoimmune diseases, such as: rheumatoid arthritis, type 1
CC diabetes, multiple sclerosis, systemic lupus erythematosus, inflammatory
CC bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious
CC anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease,
CC myocarditis, Sjogren's disease, or primary systemic vasculitis. The
CC present DNA sequence represents a human autoimmune disease-related
CC genomic-based SNP context sequence of the invention. NOTE: The present
CC sequence is not shown in the specification, but has been retrieved from
CC the WIPO website.
XX
SQ Sequence 201 BP; 49 A; 43 C; 45 G; 63 T; 0 U; 1 Other;
Query Match 25.8%; Score 200.6; DB 13; Length 201;
Best Local Similarity 99.5%; Pred. No. 4.4e-40;
Matches 200; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Qy 102 AAATACAGCTATTCTTAGGATCCTTGGCCCTTTCATAGAGATAAGTTGTGAATCCTAA 161
|||||
```

CC	bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease, myocarditis, Sjogren's disease, or primary systemic vasculitis. The present DNA sequence represents a human autoimmune disease-related genomic-based SNP context sequence of the invention. NOTE: The present sequence is not shown in the specification, but has been retrieved from the WIPO website.
CC	Sequence 201 BP; 50 A; 35 C; 47 G; 68 T; 0 U; 1 Other;
CC	Query Match 25.3%; Score 196.6; DB 13; Length 201;
CC	Best Local Similarity 99.5%; Pred. No. 4.3e-39;
CC	Matches 196; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY	1 TCATGTAGGTAAAGCTGTGTAGAAATGTCGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 60
DB	5 TCATGTAGGTAAAGCTGTGTAGAAATGTCGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 64
QY	61 TGATAAGGCTTTTGGCATTAGATGCTGTTTGTCTTCATGAGAAATACAGCTATTCTAGG 120
DB	65 TGATAAGGCTTTTGGCATTAGATGCTGTTTGTCTTCATGAGAAATACAGCTATTCTAGG 124
QY	121 ATCCTTGAGCCTTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCCTAGACCATTTACT 180
DB	125 ATCCTTGAGCCTTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCCTAGACCATTTACT 184
QY	181 TAGATGATCTGCTCTCT 197
DB	185 TAGATGATCTGCTCTCT 201
RESULT 7	
ID	ADS39163
AD	ADS39163 standard; DNA; 201 BP.
XX	AC ADS39163;
XX	DT 16-DEC-2004 (first entry)
XX	DE Human autoimmune disease-related SNP context sequence - SEQ ID 4377.
XX	OS single nucleotide polymorphism detection; SNP detection;
KW	rheumatoid arthritis; type 1 diabetes; multiple sclerosis;
KW	systemic lupus erythematosus; inflammatory bowel disease; psoriasis;
KW	thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;
KW	glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;
KW	primary systemic vasculitis; ds.
XX	Homo sapiens.
OS	WO2004083403-A2.
PN	30-SEP-2004.
XX	18-MAR-2004; 2004WO-US008461.
XX	18-MAR-2003; 2003US-0455444P.
XX	25-APR-2003; 2003US-0465241P.
XX	(APPL-) APPLERA CORP.
XX	Cargill M, Begovich AB, Alexander HC;
XX	WPI; 2004-728480/71.
XX	New isolated nucleic acid molecule comprises at least 8 contiguous nucleotides where one of the nucleotides is a single nucleotide polymorphism (SNP), useful for diagnosing or treating autoimmune diseases, e.g. rheumatoid arthritis.
XX	Claim 16; SEQ ID NO 4377; 123pp; English.
XX	The invention comprises amino acid and coding sequences containing

CC	bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease, myocarditis, Sjogren's disease, or primary systemic vasculitis. The present DNA sequence represents a human autoimmune disease-related genomic-based SNP context sequence of the invention. NOTE: The present sequence is not shown in the specification, but has been retrieved from the WIPO website.
XX	
SQ	Sequence 201 BP; 50 A; 35 C; 47 G; 68 T; 0 U; 1 Other;
Query Match	25.3%; Score 196.6; DB 13; Length 201;
Best Local Similarity	99.5%; Pred. No. 4.3e-39;
Matches 196; Conservative 1; Mismatches 0; Indels 0; Gaps 0;	
Qy	1 TCATGTAGGTAAGACTGTGTAGAATCTCGGGTCTCGATGTTCGCGCATTTCAAGCCCTGA 60
Db	5 TCATGTAGGTAAGACTGTGTAGAATCTCGGGTCTCGATGTTCGCGCATTTCAAGCCCTGA 64
Qy	61 TGATAAGGCCTTTTGCCATTAGATGCTGTTTTGTCTTCATCGAAATACAGCTATTCTTAG 120
Db	65 TGATAAGGCCTTTTGCCATTAGATGCTGTTTTGTCTTSATGGAATACAGCTATTCTTAG 124
Qy	121 ATCCTTGAGCCTTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCTAGCACCATTTACT 180
Db	125 ATCCTTGAGCCTTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCTAGCACCATTTACT 184
Qy	181 TAGATGATCTGCTCTCT 197
Db	185 TAGATGATCTGCTCTCT 201
RESULT 7	
ID	ADS39163 standard; DNA; 201 BP.
XX	ADS39163;
AC	ADS39163;
XX	
DT	16-DEC-2004 (first entry)
XX	
DE	Human autoimmune disease-related SNP context sequence - SEQ ID 4377.
XX	
KW	single nucleotide polymorphism detection; SNP detection;
KW	rheumatoid arthritis; type 1 diabetes; multiple sclerosis;
KW	systemic lupus erythematosus; inflammatory bowel disease; psoriasis;
KW	thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;
KW	glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;
KW	primary systemic vasculitis; ds.
OS	Homo sapiens.
XX	
PN	WO2004083403-A2.
XX	
PD	30-SEP-2004.
XX	
PF	18-MAR-2004; 2004WO-US008461.
XX	
PR	18-MAR-2003; 2003US-0455444P.
XX	
PT	25-APR-2003; 2003US-0465241P.
PA	(APPL-) APPLERA CORP.
XX	
PI	Cargill M, Begovich AB, Alexander HC;
XX	
DR	WPI; 2004-728480/71.
XX	
PT	New isolated nucleic acid molecule comprises at least 8 contiguous nucleotides where one of the nucleotides is a single nucleotide polymorphism (SNP), useful for diagnosing or treating autoimmune diseases, e.g. rheumatoid arthritis.
PS	Claim 16; SEQ ID NO 4377; 123pp; English.
CC	The invention comprises amino acid and coding sequences containing

SQ	Sequence 201 BP; 48 A; 41 C; 46 G; 65 T; 0 U; 1 Other;
Query Match	25.8%; Score 200.6; DB 13; Length 201;
Best Local Similarity	99.5%; Pred. No. 4.4e-40;
Matches 200; Conservative 1; Mismatches 0; Indels 0; Gaps 0;	
Qy	125 TTGAGCCTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCTAGGACCATTTACTTAGA 184
Db	1 TTGAGCCTTTCATAAGAGATAGGTTGTGAATCCTAAGACCCTAGGACCATTTACTTAGA 60
Qy	185 TGATCTGCTCTCGGTTGCTCTCTGAAAAGTCTGCTCGTGGAGGGGTGCTGCATTTG 244
Db	61 TGAFTGCTCTCGGTTGCTCTCTGAAAAGTCTGCTCGYAGGGGTGCTGCATTTG 120
Qy	245 CCTTGCCTAAGTGGTGGCACACAACCTGCTACTGTACCTTAGGCTTAAATACCATGTGT 304
Db	121 CCTTGCCTAAGTGGTGGCACACAACCTGCTACTGTACCTTAGGCTTAAATACCATGTGT 180
Qy	305 CATCTAGATGAAGTTATATT 325
Db	181 CATCTAGATGAAGTTATATT 201
RESULT 6	
ID	ADS39169 standard; DNA; 201 BP.
XX	ADS39169;
AC	ADS39169;
XX	
DT	16-DEC-2004 (first entry)
XX	
DE	Human autoimmune disease-related SNP context sequence - SEQ ID 4383.
XX	
KW	single nucleotide polymorphism detection; SNP detection;
KW	rheumatoid arthritis; type 1 diabetes; multiple sclerosis;
KW	systemic lupus erythematosus; inflammatory bowel disease; psoriasis;
KW	thyroiditis; celiac disease; pernicious anaemia; asthma; vitiligo;
KW	glomerulonephritis; Grave's disease; myocarditis; Sjogren's disease;
KW	primary systemic vasculitis; ds.
OS	Homo sapiens.
XX	
PN	WO2004083403-A2.
XX	
PD	30-SEP-2004.
XX	
PF	18-MAR-2004; 2004WO-US008461.
XX	
PR	18-MAR-2003; 2003US-0455444P.
XX	
PT	25-APR-2003; 2003US-0465241P.
PA	(APPL-) APPLERA CORP.
XX	
PI	Cargill M, Begovich AB, Alexander HC;
XX	
DR	WPI; 2004-728480/71.
XX	
PT	New isolated nucleic acid molecule comprises at least 8 contiguous nucleotides where one of the nucleotides is a single nucleotide polymorphism (SNP), useful for diagnosing or treating autoimmune diseases, e.g. rheumatoid arthritis.
PS	Claim 16; SEQ ID NO 4383; 123pp; English.
CC	The invention comprises amino acid and coding sequences containing
CC	genetic polymorphisms associated with an altered risk of developing an autoimmune disease (e.g. rheumatoid arthritis). The invention further
CC	comprises a method of identifying an individual that has an altered risk
CC	of developing an autoimmune disease, comprising detecting a single
CC	nucleotide polymorphism (SNP) in a nucleic acid of the invention. The DNA
CC	and protein sequences of the invention are useful for diagnosing and
CC	treating autoimmune diseases, such as: rheumatoid arthritis, type 1
CC	diabetes, multiple sclerosis, systemic lupus erythematosus, inflammatory

CC genetic polymorphisms associated with an altered risk of developing an
CC autoimmune disease (e.g. rheumatoid arthritis). The invention further
CC comprises a method of identifying an individual that has an altered risk
CC of developing an autoimmune disease, comprising detecting a single
CC nucleotide polymorphism (SNP) in a nucleic acid of the invention. The DNA
CC and protein sequences of the invention are useful for diagnosing and
CC treating autoimmune diseases, such as: rheumatoid arthritis, type 1
CC diabetes, multiple sclerosis, systemic lupus erythematosus, inflammatory
CC bowel diseases, psoriasis, thyroiditis, celiac disease, pernicious
CC anaemia, asthma, vitiligo, glomerulonephritis, Grave's disease,
CC myocarditis, Sjogren's disease, or primary systemic vasculitis. The
CC present DNA sequence represents a human autoimmune disease-related
CC genomic-based SNP context sequence of the invention. NOTE: The present
CC sequence is not shown in the specification, but has been retrieved from
CC the WIPO website.

XX SQ Sequence 201 BP; 62 A; 22 C; 46 G; 70 T; 0 U; 1 Other;

Query Match 25.2%; Score 195.8; DB 13; Length 201;
Best Local Similarity 98.0%; Pred. No. 6.8e-39;
Matches 197; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

Qy 411 AGTCGTGCAGTAATAGAGGTATCGTCATTCATGTGACATAAAGAGGAGGGGCTTC 470
Db 1 AGTCGTGCAGTAATAGAGGTATCGTCATTCATGTGACATAAAGAGGAGGGGCTTC 60
Qy 471 ATTCATGCTTACTGATGGAATAGGAAGTGGTGAAGTGAATTTAATAGATGTTCTTTT 530
Db 61 ATTCATGCTTACTGATGGAATAGGAAGTGGTGAAGTGAATTTAATAGATGTTCTTTT 120
Qy 531 ATGAAATAAATTTTAAAGATTGTCCAGCCCTGCATGATTTATGATGAATCATTTTGTGG 590
Db 121 ATGAAATAAATTTTAAAGATTGTCCAGCCCTGCATGATTTATGATGAATCATTTTGTGG 180
Qy 591 TCTGTTAGTACTTTTAGAGA 611
Db 181 TCTGTTAGTACTTTCTAGAGA 201

RESULT 8

ABN80072
ID ABN80072 standard; DNA; 5771 BP.

XX AC ABN80072;

XX DT 15-JUL-2002 (first entry)

XX DE Human chemically modified disease associated gene SEQ ID NO 89.

XX KW Human; development; homeobox gene; HOX; diabetes; cancer; apoptosis;
KW heart disease; epilepsy; histone deacetylation; muscular dystrophy;
KW dwarfism; single nucleotide polymorphism; SNP; cytosine methylation;
KW antidiabetic; cytosine; anticonvulsant; ds.

XX OS Homo sapiens.

XX OS Synthetic.

XX PN WO200200927-A2.

XX PD 03-JAN-2002.

XX PF 02-JUL-2001; 2001WO-EP007536.

XX PR 30-JUN-2000; 2000DE-01032529.

XX PR 01-SEP-2000; 2000DE-01043826.

XX PA (EPIG-) EPIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX XX WPI; 2002-130908/17.

XX FT Novel nucleic acid useful for diagnosis and therapy of diseases

PT associated with development genes such as diabetes, comprises a sequence
PT of a segment of chemically pretreated DNA of genes associated with
PT development.

XX Claim 1; SEQ ID NO 89; 27pp; English.

XX PS The invention relates to a nucleic acid (I) comprising a sequence at
CC least 18 bases in length of a segment of chemically pretreated DNA (II)
CC of genes associated with development selected from 87 genes listed in the
CC specification such as ACCFN, ADFN, or AFDI and comprising one of 350
CC sequences (ABN9984-ABN80333) or their complements. The invention is
CC useful for the diagnosis or therapy of diseases associated with
CC development genes, in particular disease related to homeobox containing
CC genes (HOX), like diabetes, cancer, apoptosis related diseases, syndromes
CC associated with congenital heart disease, epilepsy, diseases related to
CC histone deacetylation, Curarino syndrome, diseases related with the
CC development of the brain and limb girdle muscular dystrophy and dwarfism.
CC Oligomers specific to each of the genes are useful for detecting the
CC methylation state of all CpG dinucleotides within the 350 sequences or
CC (II) and their complementary sequences, as primer oligonucleotides for
CC the amplification of the 350 sequences, (II) and/or their complements and
CC as oligomer probes for detecting the cytosine methylation state and/or
CC single nucleotide polymorphisms (SNPs). Note: The sequence data for this
CC patent did not form part of the printed specification but is based on
CC sequence information supplied to Derwent by the European Patent Office

XX SQ Sequence 5771 BP; 1877 A; 37 C; 980 G; 2877 T; 0 U; 0 Other;

Query Match 6.4%; Score 50; DB 6; Length 5771;
Best Local Similarity 48.0%; Pred. No. 0.033;
Matches 143; Conservative 0; Mismatches 155; Indels 0; Gaps 0;

Qy 309 TAGAAGAAGTATATTTTAAAGAGGATCGTTTTGGCCATGTATAAATTTCAACATTA 368
Db 1459 TAAATTTGTGTTTTATTGTGAAAAGGATAGTTTTTTTATTATTATTATTATTTT 1518
Qy 369 ACTTTCAGGGTATTAATCCTTTTAAGGTCAGTTTTTCTTAAGTCGTGAGTAATAGA 428
Db 1519 TTATTTTGTATAGTATTTTATTGTTTAAATTTTATTGTTGTAAGTTGTAGT 1578
Qy 429 GGATATCGTCATTCATGACATATAAGATGGAAGGGCTTCATTCATGTTAGTGATGA 488
Db 1579 TTTAGTGGAAATTTTGTAGGAAAAGTTTATATATTTTATTATTATAGATATTATG 1638
Qy 489 AATAGGAAGTAGGTGAAGTGATTTTAATAGATGTTTTCTTTTATGAATAATTTTAAA 548
Db 1639 ATTATGTAATTTTATTTTGTAGTATTGTTTATAGATGTTTATTTTATAGAAGAG 1698
Qy 549 GATTGTCCAGCCCTGCATGATTTATGATGAATCATTTTGTGCTGTAGTACTTTT 606
Db 1699 AATGTAAGGATAATTAATTTAATGTTATTTTATTATTGTTAGATTAATAAATTT 1756

RESULT 9

ABL32312

ID ABL32312 standard; DNA; 18683 BP.

XX AC ABL32312;

XX DT 26-MAR-2002 (first entry)

XX DE Human immune system associated gene SEQ ID NO: 285.

XX KW Human; immune system disease; cytosine methylation; antiasthmatic;
KW antiarteriosclerotic; antianemic; cytosine; cytosine; cytosine;
KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;
KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;
KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;
KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;
KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;
KW ds.

XX OS Homo sapiens.

XX WO200200928-A2.
XX 03-JAN-2002.
XX 02-JUL-2001; 2001WO-EP007537.
XX 30-JUN-2000; 2000DE-01032529.
XX 01-SEP-2000; 2000DE-01043826.
XX (EPIG-) EPIGENOMICS AG.
XX Olek A, Piepenbrock C, Berlin K;
XX WPI; 2002-130909/17.
XX Nucleic acid comprising fragment of chemically modified gene, useful for
XX diagnosis and treatment of diseases associated with abnormal cytosine
XX methylation.
XX Claim 1; SEQ ID NO 285; 32pp + Sequence Listing; German.
XX The present invention provides a number of human immune system associated
XX genes which are modified by the methylation of cytosines. The sequences
XX can be used in the diagnosis and treatment of immune system disorders,
XX including eye diseases such as retinopathy, neovascular glaucoma and
XX macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
XX diseases. The present sequence is a gene of the invention
XX
XX Sequence 18683 BP; 5439 A; 130 C; 3594 G; 9520 T; 0 U; 0 Other;
XX
XX Query Match 6.3%; Score 49.2; DB 6; Length 18683;
XX Best Local Similarity 46.5%; Pred. No. 0.072;
XX Matches 228; Conservative 0; Mismatches 258; Indels 4; Gaps 2;
XX
XX QY 290 TTAATAACCATGTCATCTAGAGTGAAGTTATATTTTAAAGAGATCGTTTTGCCATG 349
XX DB TTAATAAATAGTCGTTGATGATTTATAATAATTAATAATGATAGTATTATTATT 6414
XX
XX QY 350 TATAAATTTTCA--AACATTAACTTTCAGGGTTATTAATCCTTTTAAAGCTAGTTTTC 407
XX DB TTAATAAATAGTCGTTGATGATTTATAATAATTAATAATGATAGTATTATTATT 6414
XX
XX QY 408 TTAAGTCGTGTCAGTAATAGAGGTATCGTTCATCTGACATAAAGATGGAAGGGGC 467
XX DB ATTAAATAGTCGTTGATGATTTTATTAATAATAAATAAATTTATAGTAAATTATTT 6534
XX
XX QY 468 TTCAATTCATGTTAGTGATGGAATAAGGAAGTAGGTGAAGTGAATTTTAAATAGATGTTTCT 527
XX DB ATTGATGTTTATTTAGTGATTTATAATAATAAATAAATAAATTTATAGTAAATTATTT 6594
XX
XX QY 528 TTTATGAATAATTTTAAAGATGTCGCCGCCCTGATGATTTATGATGATCAATTTTG 587
XX DB TTTAAAAAATATTTATATATGATAGTAAATTTATGATTTTATGATTTTAAATATTTAA 6654
XX
XX QY 588 TGGTCGTGTAGTACTTTTATAGAGATAGAAAGCATTTGAGGCTCAGGGAAGC--AAACA 645
XX DB TATTAATTAATATAGTTGTTATTTATTTATTTGATTTGATGAGGGAATTTAGAGA 6714
XX
XX QY 646 TTCAGATGAATCCOATAGAGAGGTAAATTTATTTGGCGANGPACATTTTGGCAGCCT 705
XX DB ATGTTTTTAAATATGTTTATAGTTTAAAGTTTGTGAGAGAGTGGTTTATGTTGTTGA 6774
XX
XX QY 706 AGGCTGTGACATGTCACATCTGACATGTCGTATATTTGAAATCTGTCCTTT 765
XX DB TGAATTTTAAATTTTGTATTAAATTAATTTTTCGTAATATTTTAAATATTTGTTTGG 6834
XX
XX QY 766 TTATTTGTTA 775
XX DB TTTTGTGATTA 6844

RESULT 10
ABLS4333
ID ABL54333 standard; DNA; 18683 BP.
XX AC ABL54333;
XX DT 29-JUL-2002 (first entry)
XX DE Chemically treated apoptosis gene #17.
XX KW Apoptosis; HIV; Bloom syndrome; cardiopathy; neurodegenerative disorder;
XX KW Herpes simplex virus; renal ischaemia; amyotrophic lateral sclerosis;
XX KW cancer; ds.
XX OS Unidentified.
XX PN WO200177164-A2.
XX PD 18-OCT-2001.
XX PF 06-APR-2001; 2001WO-EP003969.
XX PR 06-APR-2000; 2000DE-01019058.
XX PR 07-APR-2000; 2000DE-01019173.
XX PR 30-JUN-2000; 2000DE-01032529.
XX PR 01-SEP-2000; 2000DE-01043826.
XX PA (EPIG-) EPIGENOMICS AG.
XX PI Olek A, Piepenbrock C, Berlin K;
XX WPI; 2002-017444/02.
XX Chemically modified sequences of genes associated with apoptosis are
XX useful to determine methylation patterns of genomic DNA samples for
XX diagnosis of associated diseases such as cancer.
XX Claim 1; Seq ID #33; 24pp; English.
XX This invention relates to chemically pre-treated DNA of genes associated
XX with apoptosis. The nucleic acids are used to allocate patients for
XX specific therapy for HIV infection, Bloom syndrome, cardiopathy, aging,
XX neurodegenerative disorders, Herpes simplex virus infection, renal
XX ischaemia, amyotrophic lateral sclerosis, solid tumours and cancers. This
XX nucleotide sequence represents a chemically treated apoptosis gene. Even
XX SEQ ID numbers are the complementary DNA strands to the odd SEQ ID
XX numbers. The sequence data for this patent is not represented in the
XX printed specification but is based on information supplied by the
XX European patent office
XX
XX Sequence 18683 BP; 5439 A; 130 C; 3594 G; 9520 T; 0 U; 0 Other;
XX
XX Query Match 6.3%; Score 49.2; DB 6; Length 18683;
XX Best Local Similarity 46.5%; Pred. No. 0.072;
XX Matches 228; Conservative 0; Mismatches 258; Indels 4; Gaps 2;
XX
XX QY 290 TTAATAACCATGTCATCTAGAGTGAAGTTATATTTTAAAGAGATCGTTTTGCCATG 349
XX DB TTAATAAATAGTCGTTGATGATTTATAATAATTAATAATGATAGTATTATTATT 6414
XX
XX QY 350 TATAAATTTTCA--AACATTAACTTTCAGGGTTATTAATCCTTTTAAAGCTAGTTTTC 407
XX DB TTAATAAATAGTCGTTGATGATTTATAATAATTAATAATGATAGTATTATTATT 6414
XX
XX QY 408 TTAAGTCGTGTCAGTAATAGAGGTATCGTTCATCTGACATAAAGATGGAAGGGGC 467
XX DB ATTAAATAGTCGTTGATGATTTTATTAATAATAAATAAATTTATAGTAAATTATTT 6534
XX
XX QY 468 TTCAATTCATGTTAGTGATGGAATAAGGAAGTAGGTGAAGTGAATTTTAAATAGATGTTTCT 527
XX DB ATTGATGTTTATTTAGTGATTTATAATAATAAATAAATAAATTTATAGTAAATTATTT 6594
XX
XX QY 528 TTTATGAATAATTTTAAAGATGTCGCCGCCCTGATGATTTATGATGATCAATTTTG 587
XX DB TTTAAAAAATATTTATATATGATAGTAAATTTATGATTTTATGATTTTAAATATTTAA 6654
XX
XX QY 588 TGGTCGTGTAGTACTTTTATAGAGATAGAAAGCATTTGAGGCTCAGGGAAGC--AAACA 645
XX DB TATTAATTAATATAGTTGTTATTTATTTATTTGATTTGATGAGGGAATTTAGAGA 6714
XX
XX QY 646 TTCAGATGAATCCOATAGAGAGGTAAATTTATTTGGCGANGPACATTTTGGCAGCCT 705
XX DB ATGTTTTTAAATATGTTTATAGTTTAAAGTTTGTGAGAGAGTGGTTTATGTTGTTGA 6774
XX
XX QY 706 AGGCTGTGACATGTCACATCTGACATGTCGTATATTTGAAATCTGTCCTTT 765
XX DB TGAATTTTAAATTTTGTATTAAATTAATTTTTCGTAATATTTTAAATATTTGTTTGG 6834
XX
XX QY 766 TTATTTGTTA 775
XX DB TTTTGTGATTA 6844


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XX WO200200928-A2.
XX 03-JAN-2002.
XX 02-JUL-2001; 2001WO-EP007537.
XX 30-JUN-2000; 2000DE-01032529.
XX 01-SEP-2000; 2000DE-01043826.
XX (EPIG-) EPIGENOMICS AG.
XX Olek A, Piepenbrock C, Berlin K;
XX WPI; 2002-130909/17.
XX Nucleic acid comprising fragment of chemically modified gene, useful for
XX diagnosis and treatment of diseases associated with abnormal cytosine
XX methylation.
XX Claim 1; SEQ ID NO 192; 32pp + Sequence Listing; German.
XX The present invention provides a number of human immune system associated
XX genes which are modified by the methylation of cytosines. The sequences
XX can be used in the diagnosis and treatment of immune system disorders,
XX including eye diseases such as retinopathy, neovascular glaucoma and
XX macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
XX diseases. The present sequence is a gene of the invention
XX
XX PS Sequence 11422 BP; 3913 A; 59 C; 1856 G; 5594 T; 0 U; 0 Other;
XX
XX Query Match 6.3%; Score 49; DB 6; Length 11422;
XX Best Local Similarity 46.4%; Pred. No. 0.071;
XX Matches 235; Conservative 0; Mismatches 265; Indels 7; Gaps 2;
Qy 272 TGTACTGTCACCTTAGGCTTAATTAACCATGTCATCTAGAGTGAAGTATATATTTAAAA 331
Db 6619 TTTTATTTTAAATTTTAAAAAATTTTATTTTATTTTAAAGATATATATATTTT 6678
Qy 332 AGGATCGTTTGGCATGTATATTTTCAACATTAATTTTCAGGCTTATTAATCTTT 391
Db 6679 ATGTTGTTTAAAGATTTAAAGACGTATATATTTTAAAGTAGTGTATTA-----TT 6733
Qy 392 TAAGGTCAGTTTCTTAAAGTCGTGCAGTAATAGAGGTATCGTCATTCATGTGACATA 451
Db 6734 GAAATTTATATAGTTTAAATTTTGTGTTAATATATATAGTAGAAGATAGTATATAGA 6793
Qy 452 AAAGATGGAAGGGGCTTCATTCATGTAGTGTAGTGAAGTGAAGTGAAGTGAAT 511
Db 6794 AAATAAGAAATATAATGTTTATAGATATAAGTATATGTAATTTAAAAATTTAGTTTATTA 6853
Qy 512 TTTAATAGATGTTTCTTTTATGAATATATTTTAA--AAGATTTCAGCCCTGCATGAT 569
Db 6854 TTGAATGAGATGTTTGCATGTATTTTATTTTATTTTATATTTTATATTTTATTTAA 6913
Qy 570 TTATGATGAATCATTTTGTGTCGTGTTAGTTACTTTTATAGAAATAGAAAGCATTTGAGGC 629
Db 6914 GTTATATAAATTTTAGTATTTGTTAATTTTATTTATAGAGATTTTGTAGTATATATA 6973
Qy 630 TCAGGGAAGCAACATTCAGAAATGAATCCAAATAGAGAAGGTAATTTATTTTCGGCATG 689
Db 6974 TGATGAGAAGTTATAAATTTTGTGTTAAGATAGGAAAAATAAATGAATATTTTATTA 7033
Qy 690 TACATTTTGGCAGCTAGGCTGTACATGTGTACACATCTCGAACATGTGTGTATATTG 749
Db 7034 AATTTTATAAAGAGATATATTTTGTAAATTTTGTAGTGTATATGTTTGGAGATATATTT 7093
Qy 750 AAAATCTGTCTCTTTTATTTTATTTGTTAA 776
Db 7094 ATGTTTGGGCTTATTTTATTTTGGAA 7120
```

RESULT 13
ABK31423
ID ABK31423 standard; DNA; 13123 BP.
XX
AC ABK31423;
XX
DT 23-APR-2002 (first entry)
XX
DE Signal transduction associated gene modified complementary DNA #133.
XX
KW Human; signal transduction associated gene; cytosine methylation state;
KW CpG island; signal transduction associated disease; solid tumour; cancer;
KW antitumour; cytostatic; mutant; ds.
XX
OS Homo sapiens.
OS Synthetic.
XX
FN WO200200926-A2.
XX
PD 03-JAN-2002.
XX
PF 29-JUN-2001; 2001WO-EP007472.
XX
PR 30-JUN-2000; 2000DE-01032529.
PR
PR 01-SEP-2000; 2000DE-01043826.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K;
XX
DR WPI; 2002-147896/19.
XX
PT Oligonucleotide for diagnosis and therapy of diseases associated with
PT signal transduction e.g. cancer, comprises chemically modified genomic
PT sequences of genes associated with signal transduction.
XX
PS Claim 1; SEQ ID NO 266; 24pp; English.
XX
CC The present invention relates to chemically modified DNA sequences of
CC signal transduction associated genes. The DNA sequences are chemically
CC modified using a solution of bisulphite, hydrogen sulphite or disulphite.
CC Also disclosed are oligonucleotides and/or PNA oligomers for detecting
CC the cytosine methylation state (CpG islands) of these genes, and a method
CC for the diagnosis and/or therapy of genetic and epigenetic parameters of
CC genes associated with signal transduction. The genomic DNA can be
CC obtained from cells or cellular components which contain DNA, e.g. cell
CC lines, biopsies, blood, sputum, stool, urine, cerebral-spinal fluid,
CC tissue embedded in paraffin such as tissue from eyes, intestine, kidney,
CC brain, heart, prostate, lung, breast or liver, histologic object slides,
CC and all their possible combinations. The sequences of the invention are
CC useful for the diagnosis and therapy of diseases associated with signal
CC transduction e.g. solid tumours and cancer. ABK31158-ABK31545 represent
CC chemically pretreated genomic DNA sequences of different genes associated
CC with signal transduction, or their complementary sequences. Note: The
CC sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from the
CC European Patent Office
XX
SQ Sequence 13123 BP; 3999 A; 255 C; 2547 G; 6320 T; 0 U; 2 Other;
Query Match 6.2%; Score 48.4; DB 6; Length 13123;
Best Local Similarity 49.1%; Pred. No. 0.1;
Matches 157; Conservative 0; Mismatches 161; Indels 2; Gaps 1;
Qy 284 TTAGGCTTAATTAACCATGTCATCTAGAGTATATATTTAAAAAGATCGTTT 343
Db 2473 TTATGTATAAAATTTTATTTTATTTTATTTAGTTAAATTTTATTTAAAGATTTTGT 2532
Qy 344 GCCATGTATAAAATTTTCAACATTAACCTT--TCAGGCTTATTAATCTTTTAAGCTCTAG 401
Db 2533 TTTTGTTTTAAATTTATTTATGTTATTTTATTTTATGAGCTTTAGTTTATTTGATTTT 2592

QY 402 TTTTCTTAAGTCGTGTCAGTAATAGAGGTATCGTCATTGATGACATRAAAGATGAA 461
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 Db 2593 TTTTATTTGTTTGAAGATTAATATGGGTATTAATTTTATTAAGTTTATTTAT 2652
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 QY 462 AGGGCTTCATTCATGTTAGTGATGGAATAGGAAGTAGGTGAGTGATTTTAATAGAT 521
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 Db 2653 TAAAAATATATTTATATATAAGTTGTTTATAGGTATTTATTAAGTTTAT 2712
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 QY 522 GTTTCCTTTATGAATAATTTTAAAGATTTGTCAGCCCTGCAATGATTTATGATGATC 581
 |||||
 Db 2713 TTTTCTTTTATTTATGTTGAATTAATTTGTTAGAGTTAAGATATATTTTATTTT 2772
 |||||
 QY 582 ATTTTGGTCTGTTAGTTA 601
 |||||
 Db 2773 TTTTGGTTAGTTTGTGA 2792
 |||||

RESULT 14

ABL54364
 ID ABL54364 standard; DNA; 13123 BP.

XX ABL54364;

DT 29-JUL-2002 (first entry)

XX Chemically treated apoptosis gene complementary to gene #32.

XX Apoptosis; HIV; Bloom syndrome; cardiopathy; neurodegenerative disorder;
 KW Herpes simplex virus; renal ischaemia; amyotrophic lateral sclerosis;
 KW cancer; ds.

XX Unidentified.

XX WO200177164-A2.

XX 18-OCT-2001.

XX 06-APR-2001; 2001WO-EP003969.

XX 06-APR-2000; 2000DE-01019058.

XX 07-APR-2000; 2000DE-01019173.

XX 30-JUN-2000; 2000DE-01032529.

XX 01-SEP-2000; 2000DE-01043826.

XX (EPIG-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-017444/02.

XX Chemically modified sequences of genes associated with apoptosis are
 PT useful to determine methylation patterns of genomic DNA samples for
 PT diagnosis of associated diseases such as cancer.

XX Claim 1; Seq ID #64; 24pp; English.

XX This invention relates to chemically pre-treated DNA of genes associated
 CC with apoptosis. The nucleic acids are used to allocate patients for
 CC specific therapy for HIV infection, Bloom syndrome, cardiopathy, aging,
 CC neurodegenerative disorders, Herpes simplex virus infection, renal
 CC ischaemia, amyotrophic lateral sclerosis, solid tumours and cancers. This
 CC nucleotide sequence represents a chemically treated apoptosis gene. Even
 CC SEQ ID numbers are the complementary DNA strands to the odd SEQ ID
 CC numbers. The sequence data for this patent is not represented in the
 CC printed specification but is based on information supplied by the
 CC European patent office

XX SQ Sequence 13123 BP; 3999 A; 255 C; 2547 G; 6320 T; 0 U; 2 Other;

Query March 6.2%; Score 48.4; DB 6; Length 13123;

Best Local Similarity 49.1%; Pred. No. 0.1;

Matches 157; Conservative 0; Mismatches 161; Indels 2; Gaps 1;

QY 284 TTAGCGTTAATAACCATGTCATCTAGATTAAGTTATATTTTAAAAAGGATCGTTT 343
 |||||
 Db 2473 TTATGTATAAAATTTTATTTTAAATTTAGTTTAAATTTTATTTTAAAGTATTTTGT 2532
 |||||
 QY 344 GCCATGTATAAAATTTTCAAAACATTTAACTT--TCAGGCTTATTAATCCCTTTTAAAGGTCTAG 401
 |||||
 Db 2533 TTTTGTTTTAAATATTTATGTAATTTTATTTTATTTTATGAGGTGTTTATTTTATTT 2592
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 QY 402 TTTTCTTAAGTCGTGTCAGTAAATAGAGGTATCGTCATTCATGTCAGATAAAGATGAA 461
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 Db 2593 TTTTATTTGTTTGAAGATTAATATGCGGTATTAATTTTATTTTATTTTATTTAT 2652
 |||||
 QY 462 AGGGCTTCATTCATGTTAGTGATGGAATAGGAAGTAGGTGAGTGATTTTAAATAGAT 521
 |||||
 Db 2653 TAAAAATATATTTATATTAAGTTGTTTATAGGTATTTTATTAAGTTTAT 2712
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 QY 522 GTTTCCTTTATGAATAATTTTAAAGATTTGTCAGCCCTGCAATGATTTATGATGATC 581
 |||||
 Db 2713 TTTTCTTTTATTTATGTTGAATTAATTTGTTAGAGTTTATTTTATTTTATTTT 2772
 |||||
 QY 582 ATTTTGGTCTGTTAGTTA 601
 |||||
 Db 2773 TTTTGGTTAGTTTGTGA 2792
 |||||

RESULT 15

ABK31322
 ID ABK31322 standard; DNA; 6880 BP.

XX ABK31322;

XX 23-APR-2002 (first entry)

XX Signal transduction associated gene modified DNA #83.

XX Human; signal transduction associated gene; cytosine methylation state;
 KW CpG island; signal transduction associated disease; solid tumour; cancer;
 KW antitumour; cytostatic; mutant; ds.

XX Homo sapiens.

XX Synthetic.

XX WO200200926-A2.

XX 03-JAN-2002.

XX 29-JUN-2001; 2001WO-BP007472.

XX 30-JUN-2000; 2000DE-01032529.

XX 01-SEP-2000; 2000DE-01043826.

XX (EPIG-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-147896/19.

XX Oligonucleotide for diagnosis and therapy of diseases associated with
 PT signal transduction e.g. cancer, comprises chemically modified genomic
 PT sequences of genes associated with signal transduction.

XX Claim 1; SEQ ID NO 165; 24pp; English.

XX The present invention relates to chemically modified DNA sequences of
 CC signal transduction associated genes. The DNA sequences are chemically
 CC modified using a solution of bisulphite, hydrogen sulphite or disulphite.
 CC Also disclosed are oligonucleotides and/or PNA oligomers for detecting
 CC the cytosine methylation state (CpG islands) of these genes, and a method
 CC for the diagnosis and/or therapy of genetic and epigenetic parameters of
 CC genes associated with signal transduction. The genomic DNA can be
 CC obtained from cells or cellular components which contain DNA, e.g. cell
 CC lines, biopsies, blood, sputum, stool, urine, cerebral-spinal fluid,
 CC tissue embedded in paraffin such as tissue from eyes, intestine, kidney,

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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 15:16:42 ; Search time 197.983 Seconds
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Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 3: /cgm2_6/ptodata/1/ina/6A_COMB.seq.*
- 4: /cgm2_6/ptodata/1/ina/6B_COMB.seq.*
- 5: /cgm2_6/ptodata/1/ina/PTCUS_COMB.seq.*
- 6: /cgm2_6/ptodata/1/ina/backfiles.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	45.4	5.8	1743	4	US-09-248-796A-11015
C 2	44.4	5.7	54576	4	US-09-949-016-15954
C 3	44.4	5.7	54576	4	US-09-949-016-15955
C 4	41.8	5.4	231129	4	US-09-949-016-16110
C 5	41.8	5.4	266293	4	US-09-949-016-11934
C 6	41.4	5.3	601	4	US-09-949-016-156535
C 7	40.4	5.2	1141	4	US-09-806-708B-22
C 8	40.4	5.2	360470	4	US-09-949-016-13173
C 9	39.8	5.1	175236	4	US-09-949-016-14353
C 10	39.8	5.1	298336	4	US-09-949-016-16600
C 11	39.6	5.0	1141	4	US-09-806-708B-22
C 12	39	5.0	181429	4	US-09-949-016-142482
C 13	39	5.0	181429	4	US-09-949-016-12372
C 14	39	5.0	181430	4	US-09-949-016-15772
C 15	38.4	4.9	1783	4	US-09-679-409-9
C 16	38.4	4.9	1898	4	US-09-679-409-8
C 17	38.4	4.9	1923	4	US-09-679-409-12
C 18	38.4	4.9	1981	4	US-09-679-409-6
C 19	38.4	4.9	1992	4	US-09-679-409-7
C 20	38.4	4.9	2075	4	US-09-679-409-11
C 21	38.4	4.9	2132	4	US-09-679-409-5
C 22	38.4	4.9	2215	4	US-09-679-409-4
C 23	38.4	4.9	2226	4	US-09-679-409-2
C 24	38.4	4.9	2308	4	US-09-679-409-13
C 25	38.4	4.9	2309	4	US-09-679-409-10
C 26	38.4	4.9	2431	4	US-09-679-409-3
C 27	38.4	4.9	2612	4	US-09-679-409-15

C 28	38.4	4.9	2706	4	US-09-679-409-14	Sequence 14, Appl
C 29	38.4	4.9	319608	4	US-09-539-333D-1	Sequence 1, Appl
C 30	38.4	4.9	319608	4	US-09-679-409-1	Sequence 1, Appl
C 31	38.2	4.9	601	4	US-09-949-016-18854	Sequence 18854, A
C 32	38.2	4.9	601	4	US-09-949-016-56719	Sequence 56719, A
C 33	38.2	4.9	832	4	US-09-621-976-2813	Sequence 2813, Ap
C 34	38.2	4.9	4291	2	US-08-417-210A-80	Sequence 80, Appl
C 35	38.2	4.9	4291	4	US-09-136-159A-80	Sequence 80, Appl
C 36	38.2	4.9	4857	2	US-08-566-398-16	Sequence 16, Appl
C 37	38.2	4.9	6628	3	US-08-815-809-3	Sequence 3, Appl
C 38	38.2	4.9	6649	2	US-08-816-155B-5	Sequence 5, Appl
C 39	38.2	4.9	6649	3	US-09-079-587-5	Sequence 5, Appl
C 40	38.2	4.9	7091	2	US-08-658-665-40	Sequence 40, Appl
C 41	38.2	4.9	7091	3	US-08-796-101-4	Sequence 4, Appl
C 42	38.2	4.9	7091	3	US-09-085-273-40	Sequence 40, Appl
C 43	38.2	4.9	7091	4	US-09-916-963-40	Sequence 40, Appl
C 44	38.2	4.9	7351	1	US-08-224-391-83	Sequence 83, Appl
C 45	38.2	4.9	7351	1	US-08-484-304-83	Sequence 83, Appl

ALIGNMENTS

RESULT 1

US-09-248-796A-11015/c
; Sequence 11015, Application US/09248796A
; Patent No. 6747137
; GENERAL INFORMATION:
; APPLICANT: Keith Weinstein et al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO CANDIDA ALBICA
; FILE REFERENCE: 107196.132
; CURRENT APPLICATION NUMBER: US/09/248.796A
; CURRENT FILING DATE: 1999-02-12
; PRIOR APPLICATION NUMBER: US 60/074,725
; PRIOR FILING DATE: 1998-02-13
; PRIOR APPLICATION NUMBER: US 60/096,409
; PRIOR FILING DATE: 1998-08-13
; NUMBER OF SEQ ID NOS: 28208
; SEQ ID NO 11015
; LENGTH: 1743
; TYPE: DNA
; ORGANISM: Candida albicans
; US-09-248-796A-11015

Query Match	5.8%	Score 45.4;	DB 4;	Length 1743;
Best Local Similarity	48.6%	Pred. No. 0.028;		
Matches 124;	Conservative 0;	Mismatches 131;	Indels 0;	Gaps 0;
Qy	492	AGGAAAGTAGGTGAAGTGAATTTTAAATAGATGTTCTTTTATGAATAATATTTTAAAGAT	551	
Db	1431	AGTTAAATATTTAAATTTTGTAAATAATTTGAAATTTTCAATTAATATTAATAAAAAAT	1372	
Qy	552	TGTCACCCCTGCATGATTTATGATGATCATTTTGTGTCGTAGTTACTTTTAGAGA	611	
Db	1371	TTTGAATAATGATTTAATAATTTTCGTTTAAATTAATGATGATTTTATGATA	1312	
Qy	612	ATAGAAAGCATTTAGGCTCAGGGAAGCAAAATTCAGATGAATCCAATAGAGAAGG	671	
Db	1311	TAATAATGCAATAATTTGGTAATATATTTGAATTTTTCAGATTAATCGTTCACCGAAC	1252	
Qy	672	TAAATTTATTTGGCATGTACATTTTGGCAGCCCTAGGCTGTGTACATGTACACATTTCT	731	
Db	1251	TAAATAACCTTGATAAAATTTTTTTTTTAAATAACAATTTTCATAAAAAATTAGTCTTAATCCG	1192	
Qy	732	GAACATGTGTGTATA	746	
Db	1191	TGAGGTGGTTGAGA	1177	
US-09-949-016-15954				
Sequence 15954, Application US/09949016				

RESULT 2

```
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15954
; LENGTH: 54576
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(54576)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15954

Query Match          5.7%; Score 44.4; DB 4; Length 54576;
Best Local Similarity 50.0%; Pred. No. 0.16;
Matches 137; Conservative 0; Mismatches 136; Indels 1; Gaps 1;

Qy 271 CTGTACTGTACCTTACCTTAAACCATGTGTCATCTAGAAATGAAAGTTATATTTTAAA 330
Db 29158 CTGTAGTGTCTACTTGTGACTAAGTGGTAGACATTTGACATGAAACATATGGAAC 29217

Qy 331 AAGGATCGTTTTTGGCCATGTATAAATTTTCAACATTAACCTTTTCAGGGTTATTAATCCTT 390
Db 29218 TAATGTTATTTCGTCAGAAAGAAATTTGTTCTTAC-GGATGGTATAGGGTTAGTACATCTT 29276

Qy 391 TTAAGGTCTAGTTTTTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATCATGTGACAT 450
Db 29277 GTAAAGGCTGTTCTATTTCTGGTTTGGCCCTATCATGAGGCACAGCCCTTCAGGGGTCT 29336

Qy 451 AAAAGATGGAAGGGGCTTCATTCATGTTAGTGAAGAAATAGGAAATAGGTAAGTGAAGTGA 510
Db 29337 CAATGGAAGCCTGAAATTCATTTCTCTCTAGCACTATCAAACTACTGGATTGT 29396

Qy 511 TTTTAATAGATGTTCTTTTATGAAATAATTTTT 544
Db 29397 AGTTAAATTTTATAGCCCTTTTAGGAACATATTTTCT 29430

RESULT 3
US-09-949-016-15955
; Sequence 15955, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15955
; LENGTH: 54576
; TYPE: DNA

Query Match          5.4%; Score 41.8; DB 4; Length 231129;
Best Local Similarity 45.5%; Pred. No. 1.3;
Matches 148; Conservative 0; Mismatches 177; Indels 0; Gaps 0;

Qy 285 TAGGCTTAATAACCATGTGTCATCTAGAAATGAAAGTTATATTTTAAAAGGATCGTTTTTG 344
Db 72576 TATATTTTATATATAATAAATATATATTTTATATATAATAATAATAATAATAATAATTT 72635

Qy 345 CCATGTATAAATTTTCAACATTAACCTTCAGGGTTATTAATCCTTTTAAGGCTAGTTT 404
Db 72636 TTATTTATATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATA 72695

Qy 405 TTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATTCATGTGACATAAAGATGGAAGG 464
Db 72696 TATATATATTTATATATATATATATAATAATAATAATAATAATAATAATAATAATAATAATA 72755
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; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(54576)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15955

Query Match          5.7%; Score 44.4; DB 4; Length 54576;
Best Local Similarity 50.0%; Pred. No. 0.16;
Matches 137; Conservative 0; Mismatches 136; Indels 1; Gaps 1;

Qy 271 CTGTACTGTACCTTACCTTAAACCATGTGTCATCTAGAAATGAAAGTTATATTTTAAA 330
Db 29158 CTGTAGTGTCTACTTGTGACTAAGTGGTAGACATTTGACATGAAACATATGGAAC 29217

Qy 331 AAGGATCGTTTTTGGCCATGTATAAATTTTCAACATTAACCTTTTCAGGGTTATTAATCCTT 390
Db 29218 TAATGTTATTTCGTCAGAAAGAAATTTGTTCTTAC-GGATGGTATAGGGTTAGTACATCTT 29276

Qy 391 TTAAGGTCTAGTTTTTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATCATGTGACAT 450
Db 29277 GTAAAGGCTGTTCTATTTCTGGTTTGGCCCTATCATGAGGCACAGCCCTTCAGGGGTCT 29336

Qy 451 AAAAGATGGAAGGGGCTTCATTCATGTTAGTGAAGAAATAGGAAATAGGTAAGTGAAGTGA 510
Db 29337 CAATGGAAGCCTGAAATTCATTTCTCTCTAGCACTATCAAACTACTGGATTGT 29396

Qy 511 TTTTAATAGATGTTCTTTTATGAAATAATTTTT 544
Db 29397 AGTTAAATTTTATAGCCCTTTTAGGAACATATTTTCT 29430

RESULT 4
US-09-949-016-16110
; Sequence 16110, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16110
; LENGTH: 231129
; TYPE: DNA
; ORGANISM: Human
; OTHER INFORMATION:
US-09-949-016-16110

Query Match          5.4%; Score 41.8; DB 4; Length 231129;
Best Local Similarity 45.5%; Pred. No. 1.3;
Matches 148; Conservative 0; Mismatches 177; Indels 0; Gaps 0;

Qy 285 TAGGCTTAATAACCATGTGTCATCTAGAAATGAAAGTTATATTTTAAAAGGATCGTTTTTG 344
Db 72576 TATATTTTATATATAATAAATATATATTTTATATATAATAATAATAATAATAATAATTT 72635

Qy 345 CCATGTATAAATTTTCAACATTAACCTTCAGGGTTATTAATCCTTTTAAGGCTAGTTT 404
Db 72636 TTATTTATATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATA 72695

Qy 405 TTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATTCATGTGACATAAAGATGGAAGG 464
Db 72696 TATATATATTTATATATATATATATAATAATAATAATAATAATAATAATAATAATAATAATA 72755
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QY 465 GCCTTCATTCAGTTAGTGATGGAATAGGAAGTAGTGAGTGAATTTTAATAGATGTT 524
Db 72756 TATATATGTAATATATATTTATGTTATATATATATATATATATATATATATATAT 72815
QY 525 TCCTTTATGAATAATTTTAAAGATGTCAGCCCTGCATGATTTATGATGATCAAT 584
Db 72816 ATATACTTAAT 72875
QY 585 TTGTGGTCTGTTAGTTACTTTTGA 609
Db 72876 ATATATTTATATATAATATATATA 72900

RESULT 5

US-09-949-016-11934
; Sequence 11934, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11934
; LENGTH: 266293
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11934

Query Match 5.4%; Score 41.8; DB 4; Length 266293;
Best Local Similarity 45.5%; Pred. No. 1.3;
Matches 148; Conservative 0; Mismatches 177; Indels 0; Gaps 0;

QY 285 TAGGCTTAATACCATGTCATCTAGATGAGTATATTTTAAAAAGGATCGTTTTTG 344
Db 59740 TATATTTTATATATAAAATATATATTTTATATATATATATATATATATATTT 59799
QY 345 CCATGTATAAATTTTCAACATTAATCTTCAGGGTTATTAATCCCTTTAAGTCTAGTTT 404
Db 59800 TTATTTATATAATATATATTTTATATATATATATATATATATATATATATATAT 59859
QY 405 TTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATTCATGTGCACATAAAAGATGGAAGG 464
Db 59860 TAT 59919
QY 465 GCCTTCATTCAGTTAGTGATGGAATAGGAAGTAGTGAGTGAATTTTAATAGATGTT 524
Db 59920 TATATATGTAATATATTTATGTTATATATATATATATATATATATATATATATAT 59979
QY 525 TCCTTTATGAATAATTTTAAAGATGTCAGCCCTGCATGATTTATGATGATCAAT 584
Db 59980 ATATACTTAAT 60039
QY 585 TTGTGGTCTGTTAGTTACTTTTGA 609
Db 60040 ATATATTTATATATAATATATATA 60064

RESULT 6

US-09-949-016-156535/c
; Sequence 156535, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

Query Match

5.2%; Score 40.4; DB 4; Length 1141;

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 156535
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-156535

Query Match 5.3%; Score 41.4; DB 4; Length 601;
Best Local Similarity 45.2%; Pred. No. 0.23;
Matches 147; Conservative 1; Mismatches 177; Indels 0; Gaps 0;

QY 285 TAGGCTTAATACCATGTCATCTAGATGAGTATATTTTAAAAAGGATCGTTTTTG 344
Db 438 TATATTTTATATATAAAATATATATTTTATATATATATATATATATATATTT 379
QY 345 CCATGTATAAATTTTCAACATTAATCTTCAGGGTTATTAATCCCTTTAAGTCTAGTTT 404
Db 378 TTATTTATATAATATATATATTTTATATATATATATATATATATATATATATAT 319
QY 405 TTCTTAAGTCTGTGCAGTAATAGAGGTATCGTCAATTCATGTGCACATAAAAGATGGAAGG 464
Db 318 TATATATATTTAT 259
QY 465 GCCTTCATTCAGTTAGTGATGGAATAGGAAGTAGTGAGTGAATTTTAATAGATGTT 524
Db 258 TATATATGTAATATATATTTTATGTTATATATATATATATATATATATATATATAT 199
QY 525 TCCTTTATGAATAATTTTAAAGATGTCAGCCCTGCATGATTTATGATGATCAAT 584
Db 198 ATATACTTAAT 139
QY 585 TTGTGGTCTGTTAGTTACTTTTGA 609
Db 138 ATATATTTATATATAATATATATA 114

RESULT 7

US-09-806-708B-22/c
; Sequence 22, Application US/09806708B
; Patent No. 6784342
; GENERAL INFORMATION:
; APPLICANT: The University of British Columbia

; TITLE OF INVENTION: Regulation of Embryonic Transcription in Plants
; FILE REFERENCE: 4810-58741
; CURRENT APPLICATION NUMBER: US/09/806,708B
; PRIOR FILING DATE: 2001-04-03
; PRIOR APPLICATION NUMBER: US 60/147,133
; PRIOR FILING DATE: 1999-08-04
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: Patent in version 3.0
; SEQ ID NO 22
; LENGTH: 1141
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; NAME/KEY: promoter
; LOCATION: (1)..(1141)
; OTHER INFORMATION: consensus sequence of A.t., L.a., and B.n. FAE1 promoters
US-09-806-708B-22

Query Match

5.2%; Score 40.4; DB 4; Length 1141;

```

Best Local Similarity 9.3%; Pred. No. 0.52;
Matches 34; Conservative 155; Mismatches 178; Indels 0; Gaps 0;

Qy 316 AAGTTATATTTAAAGAGCATGTTTGGCCATGTATAAATTTTCAACATTAACCTTCA 375
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 695 WNNWTDARTNNTTVRRHRWNTNKTWYSTTTRRHHTGATNNNNNNNNNNNNNSCC 636

Qy 376 GGGTTATTAATCTTTTAAAGGCTAGTTCTTAAAGTCGTGCAGTAATAGAGGTATCG 435
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 635 TCTRMWTRWTKMGDMTVRKVKVRDITCTVYDWAWSVWVYANWRCRDVTTYRNT 576

Qy 436 TCATTTCATGTGCATATAAGAGTGAAGGGCTTCATTCATGTTAGTGATGAAATAGGA 495
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 575 YCKSYAHSYWYWSNNAWYRYSARNWSMARWTTTRNNWWSGBVWRWRWAGTWWWRHWN 516

Qy 496 AAGTAGGTGAAGTGATTTTAATAGATGTTCTTTTATGAAATAATTTTAAAGATTGTC 555
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 515 NNTDTRYWWWKRWARBTTTYYVDSMCNKMWRGNWRNWKMWAAANNIDAGAMDHTYWM 456

Qy 556 CAGCCCTCATGATTATATAGTAATCATTTTGTGTCGTAGTTACTTTTATAGAGATAG 615
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 455 GNNTWWRRAWKMMWMCRRAYCCNNNNNRACVWHKHQWRWTWKYMKWKAACNNNNBKA 396

Qy 616 AAAGCATTGTAGGCTCAGGGAAGCAAAATTCAGAATCAAAATCCAATAGAGAAGGTAA 675
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 395 MYMEVAVWMMYSRDTINTDMMWMTSDMBWHHTVYDTMRAWNNNNNNNNWRCBCKTTSWMMW 336

Qy 676 TTTATTT 682
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 335 MDHMNTH 329
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

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RESULT 8
US-09-949-016-13173
; Sequence 13173, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,769
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13173
; LENGTH: 360470
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13173

```

	Query Match	5.2%	Score 40.4;	DB 4;	Length 360470;
	Best Local Similarity	46.5%;	Prod. No. 3.5;		
	Matches 131;	Conservative	0;	Mismatches 151;	Indels 0; Gaps 0;
Qy	292	AATAACCATGTCATCTAGAAATGAAGTTATATTTTAAAAAGGATCGTTTTTGGCCATGTA	351		
Db	176767	AAAGTGCACCTATAGATACAAAACCTTTAAATTAATACAAAACGTTAGTAGTGTGATTTT	176826		
Qy	352	TAAATTTTCAAAACATTAACCTTTTCAGGGTATTAAATCCTTTTAAAGTCTAGTTTTTCTTAA	411		
Db	176827	CAC TTCATCCATTTACTTTTGCTCAGTTGATTACATAGTTTGGGGTATATACATATGTGNA	176886		
Qy	412	GTCGTGGAGTAATAGAGGTTATCGTCATTCATGTGACATATAAAGATGGAAGGGGGCTTCA	471		
Db	176887	GATACACCATATPAGACACAAAACCTTTAAATTCAGAGAAAATGTTAGACATCTAACATTTATG	176946		

Qy	472	TTCAAGTTAGTAGAGAAATAGGAAAGTAGGTGAATGATTTTAAATAGATCTTCTTTTA	531
Db	176947	GTTATATAAATTTATTGTCAATGAATTTATTGTTATGTAATTTATTTTCTCTTA	177006
Qy	532	TGAATAATATTTTAAAGATTGTCCAGCCCTGCGCATGATTTAT	573
Db	177007	TTTAGTCATTGTGATAAATATTGACGTACCTCCTTAAGCCTTAT	177048

RESULT 9
 US-09-949-016-14353
 ; Sequence 14353, Application US/09949016
 ; Patent No. 6812339
 ; GENERAL INFORMATION:
 ; APPLICANT: VENTER, J. Craig et al.
 ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 ; WITH OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CL001307
 ; CURRENT APPLICATION NUMBER: US/09/949,016
 ; CURRENT FILING DATE: 2000-04-14
 ; PRIOR APPLICATION NUMBER: 60/241,755
 ; PRIOR FILING DATE: 2000-10-20
 ; PRIOR APPLICATION NUMBER: 60/237,768
 ; PRIOR FILING DATE: 2000-10-03
 ; PRIOR APPLICATION NUMBER: 60/231,498
 ; PRIOR FILING DATE: 2000-09-08
 ; NUMBER OF SEQ ID NOS: 207012
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 14353
 ; LENGTH: 175236
 ; TYPE: DNA
 ; ORGANISM: Human
 US-09-949-016-14353

Query Match	5.1%;	Score 39.8;	DB 4;	Length 175236;
Best Local Similarity	48.1%;	Pred. No. 4;		
Matches 113;	Conservative 0;	Mismatches 122;	Indels 0;	Gaps 0;
Qy	146	AGGTTGTGAATCCTTAAGACCCTAGGACCATTTACTTATAGATGATCTGCTCTCTCGTTCGTC	205	
Db	107752	AGCTCTGTCTGATATTACTGTCTGCAGGCTTTATTTTGATTAGCAATTTCTTAGTAGTA	107811	
Qy	206	CTCTGAAAAGTCCTGCTTCGTGAGGGGTGCTGCAATTTGCCCTTAAGTGGTGTGGCA	265	
Db	107812	CAATTTTGTCTTTTCAGTCTTTCTATGTGTGTGTGTTTGTGTGACTCTTTAGAAAGATGA	107871	
Qy	266	CACAACCTGTAATGTCACCTTAAGGCTTAATAACCAATGTGCATCTAGAAATGAAGTTATATT	325	
Db	107872	TATAGCTAGATTGTTTAAAAATTCATCTATTATTTATTTAGTATGTTGAATTTTATCCCAATT	107931	
Qy	326	TTAAAAAGGATCGTTTTTGGCATGTATATAATTTTCAAACATAACTTTCAGGGTT	380	
Db	107932	ATATAGATGATCAATTAAGTATCTATTTTAGACTTATTAATCTGATTTTGTGGTT	107986	

RESULT 10
US-09-949-016-16600/c
; Sequence 16600, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FastSeq for Windows Version 4.0
 SEQ ID NO 16600
 LENGTH: 298336
 TYPE: DNA
 ORGANISM: Human
 FEATURE:
 NAME/KEY: misc_feature
 LOCATION: (1)...(298336)
 OTHER INFORMATION: n = A,T,C or G
 US-09-949-016-16600

Query Match 5.1%; Score 39.8; DB 4; Length 298336;
 Best Local Similarity 47.7%; Pred. No. 4.7; Mismatches 0; Gaps 0;
 Matches 116; Conservative 0; Indels 127; Indels 0; Gaps 0;

QY 528 TTTAAGTAATATTTTAAAGCATGTCAGCCCTGCATGATTTATGATGATCAATTTTG 587
 DB 273892 TATATATATACAGTATAGATATATAGTTATATATATATACAGTATAGATATA 273833

QY 588 TGGCTGTAGTACTTTTATAGAGATAGAAAGCATTTGAGGCTCAGGAAAGCAACATT 647
 DB 273832 TAGTTAGTTATATACAGCTATAGATATATAGTTATATATACAGTATAGATATA 273773

QY 648 CAGATGAATCCATAGAGAGGTAAATTTATTTTGGGCAATGATCAATTTTGGCAGCCTAG 707
 DB 273772 TATATATATACAGTATAGATATATAGTTATATATATATATACAGTATAGATATA 273713

QY 708 GCTGTGTACATGTTACATCTCAACATGTTGTATATTTGAAATCTTGTCTCTTTT 767
 DB 273712 GTTATATACAGTATAGATATATAGTTATATATACAGTATAGATATATATATA 273653

QY 768 TAT 770
 DB 273652 TAT 273650

RESULT 11
 US-09-806-708B-22
 ; Sequence 22, Application US/09806708B
 ; Patent No. 6784342
 ; GENERAL INFORMATION:
 ; APPLICANT: The University of British Columbia
 ; TITLE OF INVENTION: Regulation of Embryonic Transcription in Plants
 ; FILE REFERENCE: 4810-58741
 ; CURRENT APPLICATION NUMBER: US/09/806,708B
 ; PRIOR FILING DATE: 2001-04-03
 ; PRIOR APPLICATION NUMBER: US 60/147,133
 ; PRIOR FILING DATE: 1999-08-04
 ; NUMBER OF SEQ ID NOS: 23
 ; SOFTWARE: PatentIn version 3.0
 ; SEQ ID NO 22
 ; LENGTH: 1141
 ; TYPE: DNA
 ; ORGANISM: Artificial sequence
 ; FEATURE:
 ; NAME/KEY: promoter
 ; LOCATION: (1)..(1141)
 ; OTHER INFORMATION: consensus sequence of A.t., L.a., and B.n. FAE1 promoters
 US-09-806-708B-22

Query Match 5.1%; Score 39.6; DB 4; Length 1141;
 Best Local Similarity 8.8%; Pred. No. 0.85;
 Matches 55; Conservative 252; Mismatches 320; Indels 0; Gaps 0;

QY 145 AAGTTGTGAATCCCTAAGACCTTAGACCATTTACTTAGATGATCTGCTCTGTTGCT 204
 DB 77 RWWGWWYKKWYBCANNSTSBRYHARRWDMKTAYBMTWKNWKTGWRRYWRWMBDT 136

QY 205 CCTCTGAAAGTCTGCTTCTGAGGGGTGCTGCTGATTTGCTGCTGCTGCTGCTGCTGCT 264
 DB 137 VDHVYVTAANNAWTTTCMDKDDKTRWWKNNATGDDDTKYHWNNGNCBIVTWV 196

QY 265 ACACAACTGTACTGTACCTTAGGCTTAAATACCATGTCATCTAGAAATGAAGTTATAT 324

DB 197 RYKTRDWSBRRMNYGMBWKNWSYDVYVYVWVWDDMKRKRVRVRVTRGRMNTWVAWB 256
 QY 325 TTTAAAGAGGATCGTTTTTGGCATGTATAAATTTTCAACATTAACCTTTTCAGGGTTATTA 384
 DB 257 TAHRRRYNNGWIBAMAYRRWTTNNNNNAKAKRAKYGWNRABVNSTCTTWKSKTKV 316

QY 385 ATCCTTTTAAGTCTAGTCTTTTCTTAAGTCTGTGAGTAATAGAGGTATCGTCATTCATG 444
 DB 317 RTSCWANNCRAGDANKDKHKKWKSAAAGVYNNNNNNNNNTYKARHBARWVWHSWCK 376

QY 445 TGACATAAAGATGGAAGGGCTTCATTCATGTTAGTATGGAATAGGAAGTAGGTAGTG 504
 DB 377 WHANAAYSRKWTBYRKRKTVMNNNGTTWKRMMWYWKMDMDWBGTYNNNNNGRTYY 436

QY 505 AAGTGATTTTATAGATGTTCTTTTATGAATAAATTTTAAAGCATTTTCCAGCCCTGC 564
 DB 437 GWTNKKWYTKYKANNCKWRWDHKTCTHNTTWWKTYNNNNYKWSWNGSHRBA 496

QY 565 ATGATTTATGATGAATCAATTTTGTGGTCTGTAGTTACTTTTATAGAGAATAGAAAGCATG 624
 DB 497 AAVTWTWMMWRVVAHANNNNDYWKACTWYKVCCKWNNYAAWTTKSSWNTSRY 556

QY 625 TAGGCTCAGGAAAGCAACATTCAGATGAAATCCAATAGAGAAGGTAAATTTATTTGG 684
 DB 557 RWKTNNWRRSDTRSMGRANNYARABHYGYKWNTRWWSHTWBHBRAGAAHYWMBMY 616

QY 685 GCATGTACATTTTGGCAGCCTAGGCTGTGTACATGTGTACACATTTCTCAACATGTGTGA 744
 DB 617 BAKCHMKAWYKAKYAGAGSNNNNNNNNNNNNNNNNNNATCARDYYAASRYMANAKW 676

QY 745 TATTGAAATCTGTCTCTCTTTTATT 771
 DB 677 YYYKBAANNVYVTHANNWGCWNNATD 703

RESULT 12
 US-09-949-016-142482/c
 ; Sequence 142482, Application US/09949016
 ; Patent No. 6812339
 ; GENERAL INFORMATION:
 ; APPLICANT: VENTER, J. Craig et al.
 ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 ; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 ; FILE REFERENCE: CL001307
 ; CURRENT APPLICATION NUMBER: US/09/949,016
 ; CURRENT FILING DATE: 2000-04-14
 ; PRIOR APPLICATION NUMBER: 60/241,755
 ; PRIOR FILING DATE: 2000-10-20
 ; PRIOR APPLICATION NUMBER: 60/237,768
 ; PRIOR FILING DATE: 2000-10-03
 ; PRIOR APPLICATION NUMBER: 60/231,498
 ; PRIOR FILING DATE: 2000-09-08
 ; NUMBER OF SEQ ID NOS: 207012
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 142482
 ; LENGTH: 601
 ; TYPE: DNA
 ; ORGANISM: Human
 US-09-949-016-142482

Query Match 5.0%; Score 39; DB 4; Length 601;
 Best Local Similarity 46.5%; Pred. No. 1;
 Matches 126; Conservative 0; Mismatches 145; Indels 0; Gaps 0;

QY 325 TTTAAAGAGGATCGTTTTTGGCATGTATAAATTTTCAACATTAACCTTTTCAGGGTTATTA 384
 DB 406 TTTCAAACTTTTGAATATATCTTATATAAATCTATATACTATAACAAATTTTCAAA 347

QY 385 ATCCTTTTAAGTCTAGTCTTTTCTTAAGTCTGTGAGTAATAGAGGTATCGTCATTCATG 444
 DB 346 TTTCACTTCAAAATCTCTGTGGTACTCTTGAGAAAGGTGTGTGTTTTTTTTTTT 287

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Qy 445 TGACATAAAGATGGAAGGGCTTCATTGATGATGGAATAGGAAGTAGGTG 504
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Db 286 TTGGACTAAGTACCAAAATGACACATTTTGTGAAGTGGCCAAAGTAAAGTTTAA 227
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 505 AAGTGATTTTAATAGATGTTCTTTTATGAATAATTTTAAAGATGTCCAGCCCTGC 564
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 226 TTATTATAGTACACCAATTGCATGTTTGAACCAATCATGTATGAATAGTAAACTTTT 167
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 565 ATGATTTATGATGATCAATTTTGGTCTGT 595
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 166 TCTTTGTTTATAAAACCGTTTTTTTCTTT 136
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 13
US-09-949-016-12372
; Sequence 12372, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12372
; LENGTH: 181429
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12372

Query Match 5.0%; Score 39; DB 4; Length 181429;
Best Local Similarity 46.5%; Pred. No. 6.6;
Matches 126; Conservative 0; Mismatches 145; Indels 0; Gaps 0;

Qy 325 TTTAAAAAGGATCGTTTTTGCCATGATATAAATTTTCAAAACATTAACCTTCAGGGTTATTA 384
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11497 TTTACAAACTTTTGAATATCTTATATAAATACTATAAATTTTAAACAAATTTTCAA 11556
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 385 ATCCTTTTAAGGTCTAGTTTTTCTTAAGTCTGTCAGTAATAGAGGTATCGTCATTCATG 444
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11557 TTTCACTTACAAAATCTCTGTGTTACTTCTGAGAAAGGTGTGTGTTTTTTTTTTT 11616
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 445 TGACATAAAGATGGAAGGGCTTCATTGATGATGGAATAGGAAGTAGGTG 504
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11617 TTTGGACTAAGTACCAAAATGACACATTTTGTGAAGTGGCCAAAGTAAAGTTTAA 11676
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 505 AAGTGATTTTAATAGATGTTCTTTTATGAATAATTTTAAAGATGTCCAGCCCTGC 564
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11677 TTATTATAGTACCAATTGCATGTTTGAACCAATCATGTATGAATAGTAAACTTTT 11736
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 565 ATGATTTATGATGATCAATTTTGGTCTGT 595
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11737 TCTTTGTTTATAAAACCGTTTTTTTCTTT 11767
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RESULT 14
US-09-949-016-15772
; Sequence 15772, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
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; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15772
; LENGTH: 181430
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15772

Query Match 5.0%; Score 39; DB 4; Length 181430;
Best Local Similarity 46.5%; Pred. No. 6.6;
Matches 126; Conservative 0; Mismatches 145; Indels 0; Gaps 0;

Qy 325 TTTAAAAAGGATCGTTTTTGCCATGATATAAATTTTCAAAACATTAACCTTCAGGGTTATTA 384
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11497 TTTACAAACTTTTGAATATCTTATATAAATACTATAAATTTTAAACAAATTTTCAA 11556
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 385 ATCCTTTTAAGGTCTAGTTTTTCTTAAGTCTGTCAGTAATAGAGGTATCGTCATTCATG 444
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11557 TTTCACTTACAAAATCTCTGTGTTACTTCTGAGAAAGGTGTGTGTTTTTTTTTTT 11616
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 445 TGACATAAAGATGGAAGGGCTTCATTGATGATGGAATAGGAAGTAGGTG 504
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11617 TTTGGACTAAGTACCAAAATGACACATTTTGTGAAGTGGCCAAAGTAAAGTTTAA 11676
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 505 AAGTGATTTTAATAGATGTTCTTTTATGAATAATTTTAAAGATGTCCAGCCCTGC 564
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Db 11677 TTATTATAGTACCAATTGCATGTTTGAACCAATCATGTATGAATAGTAAACTTTT 11736
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Qy 565 ATGATTTATGATGATCAATTTTGGTCTGT 595
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 11737 TCTTTGTTTATAAAACCGTTTTTTTCTTT 11767
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RESULT 15
US-09-949-409-9/c
; Sequence 9, Application US/09679409
; Patent No. 655316
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Basioux, Laurent
; TITLE OF INVENTION: SCHIZOPHRENIA ASSOCIATED GENE, PROTEINS AND BIALLELIC MARKERS
; FILE REFERENCE: 53 US15 CIP
; CURRENT APPLICATION NUMBER: US/09/679,409
; CURRENT FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 09/539,333
; PRIOR FILING DATE: 2000-03-03
; PRIOR APPLICATION NUMBER: 09/416,384
; PRIOR FILING DATE: 1999-10-12
; PRIOR APPLICATION NUMBER: 60/168,088
; PRIOR FILING DATE: 1999-11-30
; NUMBER OF SEQ ID NOS: 134
; SOFTWARE: Patent.pm
; SEQ ID NO 9
; LENGTH: 1783
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..95
; NAME/KEY: CDS
; LOCATION: 96..167
; NAME/KEY: 3'UTR
; LOCATION: 168..1783
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; NAME/KEY: polyA signal
; LOCATION: 1757..1762
; NAME/KEY: allele
; LOCATION: 132
; OTHER INFORMATION: 8-130-143 : polymorphic base A or G
; NAME/KEY: allele
; LOCATION: 132
; OTHER INFORMATION: 8-126-286 : polymorphic base A or G
US-09-679-409-9

Query Match      4.9%; Score 38.4; DB 4; Length 1783;
Best Local Similarity 45.9%; Pred. No. 2.1; 196; Indels 2; Gaps 1;
Matches 168; Conservative 0; Mismatches 0;

Qy 314 TGAAGTTATATTTTAAAGGATCGTTTTGCCATGTATAAATTTTCAAAACATTAACTTT 373
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1535 TAAACAGAAATCCAAATATAGTGATATTTTCAATGCTCCTCAACTACAAATTTAATTT 1476
Qy 374 CAGGTTATTAATCCTTTTAAAGTCTAGTTTTCTTAAAGTCTGTGCAGTAATAGAGGTAT 433
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1475 TAGAGATGGCTAACCATCTATGAAGTGTATTGGATGAGCATGGTTAACAAATATTATGAAA 1416
Qy 434 CGTCATTCACTGCACATAAAGATGGAAGGGCTTCATTTCATGTTAGTGATGGAATAG 493
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1415 TAGAGTTTTTCTCACTTAAATCCAAAGAGGCCCTGAACAAGTCATTTTAGAAAAGAA 1356
Qy 494 GAAAGTAGGTGAAGTGATTTTAAATAGATGTTTCTTTATGAAATAAATTTTAAAGATTG 553
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1355 AAAAAATACTGAATAATATTTTCAAGCGTTAAATTTAGATATTAAATATAAATGAAAC 1296
Qy 554 TCCAGCCCTGCATGATTTATGATGAATCATTTTGTGCTGTGTAGTTACTTTTAGAGANT 613
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1295 TTCAACACATGPAACATATTAAGTAAC--TTTGTGTTATTTAAGCATTACCATAAAAAATTT 1238
Qy 614 AGAAGCATTGTAGGCTCAGGGAAGCAACATTCAGAAATCAATCCAAATAGAGAAGTA 673
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1237 CAAATTTATTTCTTACTTTAGATTTTAGAAGAAATTAAGTAAAGAGATCAGCCAGAGTTC 1178
Qy 674 AATTTA 679
Db ||| |||
1177 TGTGTA 1172
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Job time : 199.983 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 18:06:53 ; Search time 1005.4 Seconds
(without alignments)
6391.142 Million cell updates/sec

Title: US-09-463-542-34_COPY_368_1144

Perfect score: 777

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Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 9794790 seqs, 4134909567 residues

Total number of hits satisfying chosen parameters: 19589580

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:*

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- 2: /cgn2_6/ptodata/1/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq.*
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- 20: /cgn2_6/ptodata/1/pubpna/US10G_PUBCOMB.seq.*
- 21: /cgn2_6/ptodata/1/pubpna/US10H_PUBCOMB.seq.*
- 22: /cgn2_6/ptodata/1/pubpna/US10I_PUBCOMB.seq.*
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- 24: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq.*
- 25: /cgn2_6/ptodata/1/pubpna/US11_PUBCOMB.seq.*
- 26: /cgn2_6/ptodata/1/pubpna/US11_NEW_PUB.seq.*
- 27: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq.*
- 28: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	49.2	6.3	18683	16	US-10-311-455-285
2	49.2	6.3	18683	17	US-10-240-452-33
3	49	6.3	11422	16	US-10-311-455-192
4	49	6.3	11422	18	US-10-257-166-18
5	48.4	6.2	13123	17	US-10-240-452-64

c	6	48	6.2	595	13	US-09-925-065A-814225	Sequence 814225,
c	7	48	6.2	1693	14	US-10-027-632-256987	Sequence 256987,
c	8	48	6.2	1693	14	US-10-027-632-256988	Sequence 256988,
c	9	48	6.2	1693	18	US-10-027-632-256987	Sequence 256987,
c	10	48	6.2	1693	18	US-10-027-632-256988	Sequence 256988,
c	11	48	6.2	3673778	17	US-10-312-841-2	Sequence 2, Appl1
c	12	47	6.0	6880	18	US-10-221-613-183	Sequence 183, Appl
c	13	46.8	6.0	6179	19	US-10-221-714A-65	Sequence 65, Appl
c	14	46.6	6.0	33053	20	US-10-433-793-36	Sequence 36, Appl
c	15	46.6	6.0	3673778	17	US-10-312-841-1	Sequence 1, Appl1
c	16	46.4	6.0	50000	22	US-10-706-635-26	Sequence 26, Appl
c	17	45.2	5.8	2000	9	US-09-938-842A-5299	Sequence 5299, Ap
c	18	45.2	5.8	2000	11	US-09-938-842A-5299	Sequence 5299, Ap
c	19	45.2	5.8	5718	16	US-10-311-455-1346	Sequence 1346, Ap
c	20	45.2	5.8	5718	19	US-10-221-714A-186	Sequence 186, Ap
c	21	44.4	5.7	5886	16	US-10-311-455-2186	Sequence 2186, Ap
c	22	44.4	5.7	5992	18	US-10-221-613-167	Sequence 167, Ap
c	23	44.4	5.7	16811	16	US-10-311-455-1919	Sequence 1919, Ap
c	24	44.4	5.7	18624	16	US-10-311-455-1675	Sequence 1675, Ap
c	25	44.2	5.7	655	13	US-09-925-065A-776150	Sequence 776150,
c	26	44.2	5.7	657	13	US-09-925-065A-790118	Sequence 790118,
c	27	44.2	5.7	5937	16	US-10-240-485-95	Sequence 95, Appl
c	28	44.2	5.7	6823	16	US-10-311-455-1118	Sequence 1118, Ap
c	29	44	5.7	13584	16	US-10-311-455-588	Sequence 588, App
c	30	43.8	5.6	17918	18	US-10-221-613-381	Sequence 381, App
c	31	43.4	5.6	2000	9	US-09-938-842A-3892	Sequence 3892, Ap
c	32	43.4	5.6	2000	11	US-09-938-842A-3892	Sequence 3892, Ap
c	33	43.4	5.6	6849	19	US-10-240-589C-62	Sequence 62, Appl
c	34	43.4	5.6	10543	18	US-10-221-613-120	Sequence 120, App
c	35	43.2	5.6	6056	16	US-10-311-455-999	Sequence 999, App
c	36	43.2	5.6	7516	16	US-10-311-455-33	Sequence 33, Appl
c	37	43	5.5	6565	19	US-10-221-714A-188	Sequence 188, App
c	38	43	5.5	13321	19	US-10-221-714A-144	Sequence 144, App
c	39	42.8	5.5	6022	19	US-10-221-714A-383	Sequence 383, App
c	40	42.8	5.5	7072	18	US-10-221-613-347	Sequence 347, App
c	41	42.8	5.5	12393	16	US-10-311-455-1235	Sequence 1235, Ap
c	42	42.6	5.5	7676	16	US-10-240-485-152	Sequence 152, App
c	43	42.6	5.5	21537	16	US-10-311-455-1971	Sequence 1971, Ap
c	44	42.4	5.5	1032	18	US-10-282-122A-16418	Sequence 16418, A
c	45	42.4	5.5	7384	16	US-10-311-455-719	Sequence 719, App

ALIGNMENTS

RESULT 1

US-10-311-455-285
; Sequence 285, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPENBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Determining Cytosine Methylation
; FILE REFERENCE: 5013.1014
; CURRENT FILING DATE: 2002-12-16
; PRIOR FILING DATE: 2001-07-02
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 2424
; SEQ ID NO 285
; LENGTH: 18683
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Chemically treated genomic DNA (Homo sapiens)
US-10-311-455-285

6679	ATGTTGTTTAAAGATTTAAAGACGTATATATATTTTAAAGTAGTGTATTATA-----TT	6733
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6734	GAAATTTATATATAGTGTCTTAAATTTGCTGTTAATATATAGTGTAGAGAAAGATAGTAAATATAGA	6793
452	AAAGATGGAAGGGGCTTCATTCATGTTAGTTCGAAATAGGAAATAGTAGTGAAGTGAT	511
6794	AAATAAGAAATATAAATGTTTTAGATATAAGTATAGTATAAATAAATTTAGTTTTATTA	6853
512	TTTAATAGATGTTTTCTTTTATGAAATAAATTTTAA--AAGATTGTCAGCCCTGCGATGAT	569
6854	TTCAATGGATGTTTTTGATATGATATTTTTTTTTTAAATTTTATATATTTTAATTTATTTAA	6913
570	TTATATGATGATCATTTTGTGCTCTGTAGTACTCTTTTAGAGATAGAAAGCATTTGTAGGC	629
6914	GTTATATAAATTTTAGTATTTGGTAAATTTTTTTTATTTAGAAAGATTTTGTAGGTAGTTATATA	6973
630	TCAGGAAAGCAACATTCAGAAATGAAATCCAAATAGAGAAAGTAAATTTATTTGGGCATG	689
6974	TGATGAGAAGTTATAATTTTTTGGTTAAGATAGGAAAAATAAAATGAAATATTTTATTATA	7033
690	TACATTTTGGCAGCCTAGGCTGTGTACATGTGTACATTTCTGAACATGTGTGTATATTG	749
7034	AATTTTATAAAGAGATATATTTTGTAAATTTGTAGTGATATGTTTGTGGAGATAATATTT	7093
750	AAAATCTGTCTCTTTTTTATATGTTAA	776
7094	ATGTTTTTGGCGTTATATTTTTTTTTTGAA	7120

RESULT 5
US-10-240-452-64

```

1 PUBLICATION NO.: US20030162194A1
2 GENERAL INFORMATION:
3 APPLICANT: OLEK, Alexander
4 APPLICANT: PIEPENBROCK, Christian
5 APPLICANT: BERLIN, Kurt
6 TITLE OF INVENTION: Diagnosis of Diseases Associated with Apoptosis
7 FILE REFERENCE: 5013.1006
8 CURRENT APPLICATION NUMBER: US/10/240,452
9 CURRENT FILING DATE: 2002-10-02
10 PRIOR APPLICATION NUMBER: PCT/EP01/03969
11 PRIOR FILING DATE: 2001-04-06
12 PRIOR APPLICATION NUMBER: DE 10019058.8
13 PRIOR FILING DATE: 2000-04-06
14 PRIOR APPLICATION NUMBER: DE 10019173.8
15 PRIOR FILING DATE: 2000-04-07
16 PRIOR APPLICATION NUMBER: DE 10032529.7
17 PRIOR FILING DATE: 2000-06-30
18 PRIOR APPLICATION NUMBER: DE 10043826.1
19 PRIOR FILING DATE: 2000-09-01
20 NUMBER OF SEQ ID NOS: 78
21 SEQ ID NO 64
22 LENGTH: 13123
23 TYPE: DNA
24 ORGANISM: Artificial Sequence
25 FEATURE:
26 OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
27 FEATURE:
28 NAME/KEY: unsure
29 LOCATION: (8274.. )
30 US-10-240-452-64

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; LOCATION: ( 8274...)
US-10-240-452-64

Query Match: 6.2%; Score 48.4; DB 17; Length 13123;
Best Local Similarity 49.1%; Pred. No. 0.61;
Matches 157; Conservative 0; Mismatches 161; Indels 2; Gaps 1;

Qy 284 TTAGGCTTAATAACCATGTGTCATCTAGAAATGAAGTTATATATTTAAAAAGGATCGTTTT 343
Db 2473 TTATGTATAAAATTTTTTATTTTAAATTTAGTPTTAAATTTTTTAAAAAGTATTTTGT 2532

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, PRIOR APPLICATION NUMBER: US 60/218,006
 , PRIOR FILING DATE: 2000-07-12
 , PRIOR APPLICATION NUMBER: US 60/198,676
 , PRIOR FILING DATE: 2000-04-20
 , PRIOR APPLICATION NUMBER: US 60/193,483
 , PRIOR FILING DATE: 2000-03-29
 , PRIOR APPLICATION NUMBER: US 60/185,218
 , PRIOR FILING DATE: 2000-02-24
 , PRIOR APPLICATION NUMBER: US 60/167,363
 , PRIOR FILING DATE: 1999-11-23
 , PRIOR APPLICATION NUMBER: US 60/156,358
 , PRIOR FILING DATE: 1999-09-28
 , PRIOR APPLICATION NUMBER: US 60/146,002
 , PRIOR FILING DATE: 1999-08-09
 , NUMBER OF SEQ ID NOS: 325720

547 AAGATTGTCAGCCCGCATGATTATGATGAATCATTTTGGTCTGTAGTTACTTTT 606

Db 1672143 TAGTTTTTAAATGGTGGTGTTATTTTTTATATAATATTGTAATAAATTTTAAGATTTCGTTT 1672202
Qy 607 AGAGAATAGAAAGCATT 623
Db 1672203 TGGGACGTAATATCGTT 1672219

Search completed: November 8, 2005, 23:53:38
Job time : 1012.4 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 22:16:08 ; Search time 4014.41 Seconds
(without alignments)
7367.447 Million cell updates/sec

Title: US-09-463-542-34_COPY_368_1144

Perfect score: 777

Sequence: 1 tcattgtagtaagactgtgt.....gtctctttttttattgttaag 777

Scoring table: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*

1: gb_est1.*
2: gb_est2.*
3: gb_hic.*
4: gb_est3.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_gss1.*
9: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	101.6	13.1	376	2	BF736184
C 2	63.4	8.2	358	9	CT774457
C 3	62.4	8.0	390	1	AU234451
C 4	62.4	8.0	400	1	AU278852
C 5	58.2	7.5	1101	9	CNS0039G
C 6	54.6	7.0	919	9	CNS005RL
C 7	53.8	6.9	1101	9	CNS005EV
C 8	53.8	6.9	1101	9	CNS00LT2
C 9	53.8	6.9	1101	9	CNS016LI
C 10	52.4	6.7	1200	9	CNS016CO
C 11	52.4	6.7	1146	9	CNS021G2
C 12	51.8	6.7	1530	9	AG382019
C 13	51.6	6.6	500	1	AU088479
C 14	51.6	6.6	1201	9	CNS016BY
C 15	51.4	6.6	945	9	CNS04DOK
C 16	50.2	6.5	1190	9	CNS02ON7
C 17	49.6	6.4	1101	9	CNS00GCK
C 18	48.8	6.3	928	9	CNS00DKY
C 19	48.6	6.3	996	9	CNS00FUH
C 20	48.4	6.2	1001	9	CNS01400
C 21	48.4	6.2	1101	9	CNS0100X
C 22	48.4	6.2	1201	9	CNS0162X
C 23	48.4	6.2	1101	9	CNS001FB
C 24	47.8	6.2	842	8	BZ696583

25	47.6	6.1	979	9	CNS0161W
C 26	47.6	6.1	1315	3	CR657214
C 27	47.4	6.1	813	9	EX218284
C 28	47.4	6.1	1092	9	CNS020K7
C 29	47.4	6.1	1169	9	CNS06KHQ
C 30	47.2	6.1	1101	9	CNS000D1
C 31	47	6.0	1065	8	BZ696860
C 32	47	6.0	1094	9	CNS012FZ
C 33	47	6.0	1101	9	CNS00265
C 34	46.8	6.0	510	1	AV778356
C 35	46.8	6.0	609	9	CNS025K2
C 36	46.8	6.0	1031	9	CNS000CP2
C 37	46.8	6.0	1101	9	CNS003BD
C 38	46.8	6.0	1101	9	CNS000ES1
C 39	46.6	6.0	792	9	CG116113
C 40	46.6	6.0	870	9	CG209274
C 41	46.6	6.0	878	9	CNS0187R
C 42	46.6	6.0	894	8	BZ687506
C 43	46.6	6.0	906	9	CG209262
C 44	46.4	6.0	1896	9	CG753083
C 45	46	5.9	516	5	BM883909

ALIGNMENTS

RESULT 1
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LOCUS BF736184 376 bp mRNA linear EST 10-JAN-2001
DEFINITION PM4-KT0005-151100-001-b03 KT0005 Homo sapiens CDNA, mRNA sequence.
ACCESSION BF736184
VERSION BF736184.1 GI:12062858
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 376)
AUTHORS Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.P.,
Goldman,G.H., Carvalho,A.P., Mateukuma,A., Baia,G.S., Simpson,D.H.,
Brunstein,A., deOliveira,P.S., Bucher,P., Jongenseel,C.V.,
O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
Simpson,A.J.
TITLE Shotgun sequencing of the human transcriptome with ORF expressed
sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE 20202663
PUBMED 10737800
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the PAPESP/LICR Human Cancer Genome
Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?cl=PM4&t2=PM4-KT0005-
151100-001-b03&t3=2000-11-15&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 21
High quality sequence stop: 376.

FEATURES
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/mol_type="mRNA"
/db_xref="taxon:9606"
/dev stage="Adult"
/clone lib="KT0005"
/note="Organ: bladder tumor; Vector: puc18; Site:1: SmaI;
Site:2: SmaI; A mini-library was made by cloning products

derived from ORBSTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN

Query Match 13.1%; Score 101.6; DB 2; Length 376;
 Best Local Similarity 95.8%; Pred. No. 8.4e-14;
 Matches 115; Conservative 0; Mismatches 4; Indels 1; Gaps 1;
 Qy 1 TCATGTAGTAACTGTGTAGAGTGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 60
 Db 122 TCATGTAGTAACTGTGTAGAGTGGGTCTCGATGTGGCGCTATTCAAGCCCTGA 63
 Qy 61 TGATAAGCCTTTGGCATTAGATCTGTTTGTCTTCATCGAAATACAGCTATTCTAGG 120
 Db 62 TGATAAGCCTTTTGTCTATTAGATCTGTTTGTCTTCATCG-ATACACAGCTATTCTGAGG 4

RESULT 2

CC774457/c
 LOCUS CC774457 358 bp DNA linear GSS 27-JUN-2003
 DEFINITION CH240_95D2.TV CHORI-240 Bos taurus genomic clone CH240_95D2,
 genomic survey sequence.
 ACCESSION CC774457
 VERSION CC774457.1 GI:32326748
 KEYWORDS GSS.
 SOURCE Bos taurus (cow)
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 Bovinae; Bos.
 REFERENCE 1 (bases 1 to 358)
 AUTHORS Larkin,D.M., Everts-van der Wind,A., Rebeiz,M., Schweitzer,P.,
 Bachman,S., Green,S., Campos,E.J., Benson,L.D., Edwards,J., Liu,L.,
 Womack,J.E., de Jong,P.J. and Lewin,H.A.
 TITLE Bovine BAC end sequences from CHORI-240 library
 JOURNAL Unpublished (2003)
 COMMENT Other_GSSs: CH240_95D2.TJ
 Contact: Harris Lewin
 Department of Animal Sciences
 University of Illinois at Urbana Champaign
 1201 W. Gregory Dr., Urbana, IL 61801, USA
 Tel: 217 333 5998
 Fax: 217 244 5617
 Email: h-lewin@uiuc.edu
 Clones are derived from the bovine BAC library CHORI-240
 (<http://www.chori.org/bacpac/bovine240.htm>). For BAC library
 availability, please contact Pieter de Jong (pdejong@mail.cho.org).
 Clones may be purchased from BACPAC Resources
 (<http://www.chori.org/bacpac/ordering-information.htm>). This work
 was undertaken as part of the International Bovine BAC Mapping
 Consortium (IBBMC) by the University of Illinois at Urbana
 Champaign, USA with funds provided by grant No. AG202-34480-11828
 from USDA-CSREES and AG99-35205-8534 from USDA/NRI (Livestock
 Genome Sequencing Initiative)
 Plate: 95 row: D column: 2
 Seq primer: T7
 Class: BAC ends.

FEATURES
source

Location/Qualifiers
 1..358
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 /db_xref="taxon:9913"
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 /clone_lib="CHORI-240"
 /note="vector: pPARBAC1.3; Site.1: MboI; Site.2: MboI;
 Hereford Bull L1 Domino 99375; CHORI-240 Bovine BAC
 library (Male) produced by Pieter de Jong"

ORIGIN

Query Match 8.2%; Score 63.4; DB 9; Length 358;
 Best Local Similarity 71.5%; Pred. No. 0.00014;
 Matches 98; Conservative 0; Mismatches 36; Indels 3; Gaps 1;
 Qy 283 CTTAGGCTTAATAACCATGTGTCTAGTAAGTATATATTTTAAAGGATCGTTTT 342
 Db 140 CTTTAGGTAGCAGGCATGTGCCCTCTAGAACAAAGCTTGTTATTTTCAATAATGTCATCTT 81
 Qy 343 TGCATGTATATAATTTTCAAAACAT---TAACTTTACGGGTTATTAATCCCTTTTAAGTCT 399
 Db 80 TGCTGTGTATAAATCTTTATCAATCAACCTATCTTTTGAGGTTAGTAATCCCTTTTAAGATCT 21
 Qy 400 AGTTTTCTTTAAAGTCTG 416
 Db 20 AGTTTTCTTTAAAGTCTG 4

RESULT 3

AU234451/c
 LOCUS AU234451 390 bp mRNA linear EST 21-SEP-2001
 DEFINITION AU234451 Bovine placenta cDNA Bos taurus cDNA clone Cln153 3',
 mRNA sequence.
 ACCESSION AU234451
 VERSION AU234451.1 GI:15719669
 KEYWORDS EST.
 SOURCE Bos taurus (cow)
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 Bovinae; Bos.
 REFERENCE 1 (bases 1 to 390)
 AUTHORS Gohma,H., Lejukole,H.Y., Taniguchi,Y., Yamada,T., Akagi,S.,
 Yasue,H. and Sasaki,Y.
 TITLE Analysis of expressed sequence tags from a cDNA library of bovine
 placenta
 JOURNAL Unpublished (2001)
 COMMENT Contact: Takahisa Yamada
 Graduate School of Agriculture
 Kyoto University
 Sakyo-ku, Kitashirakawa, Kyoto, Kyoto 606-8502, Japan
 Tel: 81-75-753-6323
 Fax: 81-75-753-6340
 Email: tyamada@ikans.ikans.kais.kyoto-u.ac.j
 This clone was obtained from a 3' end cDNA library.

FEATURES
source

Location/Qualifiers
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 /organism="Bos taurus"
 /mol_type="mRNA"
 /db_xref="taxon:9913"
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 /clone_lib="Bovine placenta cDNA"

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 Best Local Similarity 69.1%; Pred. No. 0.00025;
 Matches 123; Conservative 0; Mismatches 36; Indels 19; Gaps 2;
 Qy 600 TACTTTTAGAATAGAAAGCATTAGGCTCAGGGAACAAACATTCAGATGAATC 659
 Db 299 TAGTCTTAGAACAACAAGTGTTTAAAGCTCAGGGAACAAACATTCAGATGAATC 240
 Qy 660 CAATAGAGAGTTAAATTTATTTGGGCATGTACATTTGGCAGCTAGGCTGTACATG 719
 Db 239 CAAC-----ATGTACTTCTGCTTATCATTTTGACAGATGTTGTTAGT- 192
 Qy 720 TGTACACATTCCTGAACATGTGTGTATATTGAAATCTTGTCTCTTTTATTGTTAAG 777
 Db 191 -----ATTTCGAATATGTGTATATTAAATCTCTTCTTTTCAGTTGTCAG 141

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RESULT 4
AU278852
LOCUS
DEFINITION AU278852 400 bp mRNA linear EST 02-JUL-2002
            placenta1153 3', mRNA sequence.
ACCESSION AU278852.1 GI:21682162
VERSION
KEYWORDS
SOURCE
ORGANISM Bos taurus (cow)
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
            Bovinae; Bos.
REFERENCE 1 (bases 1 to 400)
AUTHORS Oishi,M., Yamada,T., Goma,H., Lejukole,H.Y., Taniguchi,Y. and
            Sasaki,Y.
TITLE EST analysis of cloned bovine fetus and placenta
JOURNAL Unpublished (2002)
COMMENT Contact: Masahito Oishi
            Graduate School of Agriculture
            Sakyo University
            Sakyo Kitashirakawa, Kyoto, Kyoto 606-8502, Japan
            Tel: 81-75-753-6331
            Fax: 81-75-753-6340
            Email: oishi@kans.jkans.kais.kyoto-u.ac.jp.
FEATURES
            source
            Location/Qualifiers
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                /mol_type="mRNA"
                /db_xref="taxon:9913"
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                /dev_stage="60 embryonic day"
                /clone_lib="Cloned bovine placenta cdna"
ORIGIN
Query Match 8.0%; Score 62.4; DB 1; Length 400;
Best Local Similarity 69.1%; Pred. No. 0.0025;
Matches 123; Conservative 0; Mismatches 36; Indels 19; Gaps 2;
QY 600 TACTTTAGAGAAAGCAATCTAGGCTCAGGGAAGCAAAATTCAGAAATCAATC 659
Db 92 TAGTTCTAGAGAACAAAGTGTCTTTAGCTCAGGGAAGCAAAATTCAGATGAATC 151
QY 660 CAATAGAGAAGTAAATTTATTTGGCGATGACATTTTGGCAGCTAGGCTGTGTACATG 719
Db 152 CAAC-----ATGACTCTCTGCTATCCATTTTGACAGAGAGTTTGTCTAAGT- 199
QY 720 TGTACACATTCGACATGCTGTATTTGAAATCTTCTCTTTTATTTATTTAAG 777
Db 200 -----ATTTGAATATGTGTATATTAATAATCTTCTTTTTCAGTTGTTTCAG 250

RESULT 5
CNS0039G/c
LOCUS
DEFINITION CNS0039G 1101 bp DNA linear GSS 03-JUN-1999
            Drosophila melanogaster genome survey sequence TET3 end of BAC #
            BACR08K10 of RPCI-98 library from Drosophila melanogaster (fruit
            fly), genomic survey sequence.
ACCESSION AL063921
VERSION
KEYWORDS
SOURCE
ORGANISM Drosophila melanogaster (fruit fly)
            Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
            Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
            Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1101)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
            BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
            - Web : www.genoscope.cns.fr)
COMMENT Determination of this BAC-end sequence was carried out as part of a

```

collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osogawa and Aaron Mammoss in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

source

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ORIGIN

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Best Local Similarity 14.3%; Pred. No. 0.0031;
Matches 66; Conservative 223; Mismatches 172; Indels 0; Gaps 0;
QY 309 TAGAATGAAGTATATTTTAAAGGATCGTTTGGCCATGTATATAATTTCAACATTA 368
Db 1049 TWTATATWWWATWDTWWDKWWATAAKTDTATWMTAMRADWAGDRGAGKDR 990
QY 369 ACTTTCAGGGTATTAACTCTTTAAGGTCTAGTTTCTTAAAGTCTGTCAGTAATAGA 428
Db 389 DAATDADGAGRRDGGRRKDKDKDKGDDKGGKKKAAKAAKATKWDWDWDKDKW 930
QY 429 GGTATCGTCATTCATGTGACATAAAGATGAAAGGGCTTCATTCATGTAGTCATGA 488
Db 929 WDGAQRKADDDDCAGDKDDGDKGADDDTDTGDKDDDKDDDKDKDKGWTGATWAW 870
QY 489 AATAGGAAGTAGTGAAGTCAATTTTAATAGATGTTTCTTTTATGAATAATTTTAAA 548
Db 869 AATDWWMGWADADAWTWDAAADDWADDDRDWDAWAWKWDADAWGARTADRDWDG 810
QY 549 GATTGTCAGCCCTGCGATGATTATGATGAATCATTTTGTGCTGTGTAGTTACTTTAG 608
Db 809 RGGARKDRKXADDKDDAADDKDDAATWTTTDTDDDKWKTDTWTWAAADRTWD 750
QY 609 AGAATAGAAACATTTAGGCTCAGGGAAGCAAAATTCAGAAATGAATCAATAGAGA 668
Db 749 RDDDDDRDRAGTAGRKWRRTWKRKRDRTRWDDADADDTARDDRRRRGGDDGADAGK 690
QY 669 AGTAAATTTATTTGGGCATGTACATTTTGGCAGCTAGGCTGTGTACATGTGTACAT 728
Db 689 TGRKRRDRDRATWDRDADAWADAAWTTTDTDDDKRRRRKRRRRRTTAAADWD 630
QY 729 TCTGAACATGTGTATATTCGAAATCTTGTCTCTTTTAA 769
Db 629 WTWKANDWAKWDKTRADRWDRWAADTWDKADRDWAKA 589

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RESULT 6

CNS005RL/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

6

919 bp DNA linear GSS 03-JUN-1999

Drosophila melanogaster genome survey sequence T7 end of BAC #

BACR12F23 of RPCI-98 library from Drosophila melanogaster (fruit

fly), genomic survey sequence.

AL061409

GI:4943512

Drosophila melanogaster (fruit fly)

Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;


```

QY 768 TATTGTT 774
Db 491 WATAAWT 485

RESULT 8
CNS00L72
LOCUS
DEFINITION
Drosophila melanogaster genome survey sequence TET3 end of BAC:
BACR48P19 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AL078714
ACCESSION
VERSION AL078714.1 GI:5102004
KEYWORDS
SOURCE
ORGANISM
Drosophila melanogaster (fruit fly)
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Determination of this BAC-end sequence was carried out as part of a
collaboration with the Berkeley Drosophila Genome Project (BDGP).
The BDGP is constructing a physical map of the Drosophila
melanogaster genome using these BACs. For further information
please see http://www.fruitfly.org The BDGP Drosophila
melanogaster BAC library was prepared by Kazutoyo Osoegawa and
Aaron Mammosier in Pieter de Jong's laboratory in the Department of
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
NY. The library is named RPCI-98 and was constructed by partial
EcoRI digestion of Drosophila DNA provided by the BDGP from the
isogenic strain y2; cn bw sp, the same strain used for the BDGP's
P1 and EST libraries. A more detailed description of the library
and how to order individual BAC clones, the entire library, or
filters for hybridization from the BACPAC Resource Center can be
found at http://bacpac.med.buffalo.edu/drosophila\_bac.htm.

FEATURES
Location/Qualifiers
1..1101
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BACR48P19"
/clone_lib="RPCI-98"
/note="end : TET3"

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Best Local Similarity 19.7%; Pred. No. 0.035;
Matches 91; Conservative 163; Mismatches 209; Indels 0; Gaps 0;

QY 312 AATGAAGTATATTTTAAAGAGATCGTTTGGCATGTATAAATTTTCAACATAACT 371
Db 569 AAKTAAGAAWTTTATTTTAAATTTTAKTKTWAKWAWADTATTTTWTWT 628

QY 372 TTCAGGGTATTAATCCCTTTTAAGTCTAGTTTCTTAACTCTGTCAGTAATAGAGT 431
Db 629 TTTWAATTTTWTWATAWTTTTTTTAAKTATKTTTTTTTTATTAATAAATAAATTT 688

QY 432 ATCGTCATTCATGTACATAAAGATGAAAGGGCTTCATTCATGTTAGTGATGAAT 491
Db 689 TDTWAAANWTTTKKKKKAADKWKDAKKWDGAKKATTKKKDKKAWAADKKORKK 748

QY 492 AGGAAGTAGGTGAAGTGAATTTAATAGATGTTCTTTTATGAATAAATTTTAAAGAT 551
Db 749 GKKGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGG 808

QY 552 TGTCCAGCCCTGCATGATTATATGATGATCATTTTGTGTCGTGTTAGTACTTTAGAGA 611
Db 552 TGTCCAGCCCTGCATGATTATATGATGATCATTTTGTGTCGTGTTAGTACTTTAGAGA 611

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Db 809 GKKKKKAAKKKKKAADRTKTKTKWDAAAAAAAKKTKDKGKKKKKTKTKTKKKKKKKKK 868
QY 612 ATAGAAAGCATTTAGGCTCAGGGAAGCAACAACTTTCAGAAATGAAATCAAGAGAAGG 671
Db 869 GGGKDDAAAKKKKGGTGGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGG 928
QY 672 TAAATTTTGGGATGTACATTTTGGCAGCCCTAGGCTGTGTACATGTGTACATTTCT 731
Db 929 GKKGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGGKGG 988
QY 732 GAACATGTGTCTATATGAAATCTTGTCTCTTTTATTGTT 774
Db 989 KKKKKDKKKTKTKKKDAAAAAATKKKKKKKKTKTKTKTKKKKK 1031

RESULT 9
CNS016LI
LOCUS
DEFINITION
Drosophila melanogaster genome survey sequence T7 end of BAC:
BACN16D22 of DrosBAC library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AL106896
ACCESSION
VERSION AL106896.1 GI:5624374
KEYWORDS
SOURCE
ORGANISM
Drosophila melanogaster (fruit fly)
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk - This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaud at CSHF (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Payan. It has been constructed in the vector
pBelobAC11.

FEATURES
Location/Qualifiers
1..1101
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BACN16D22"
/clone_lib="DrosBAC"
/plasmid="pBelobAC11"
/note="end : T7"

ORIGIN
Query Match 6.9%; Score 53.8; DB 9; Length 1101;
Best Local Similarity 30.2%; Pred. No. 0.035;
Matches 114; Conservative 86; Mismatches 177; Indels 0; Gaps 0;

QY 297 CCATGTCTCTAGATAGTTATATTTTAAAGAGATCGTTTGGCATGTATAAAT 356
Db 725 CCCTGWRAGGGWAAWAAAAAATKATDTTTTTTTTWTWTWTWTWTWTWTWTWT 784

QY 357 TTTCAACATTAACCTTTTCAGGGTTATTAATCCCTTTTAAGGCTAGTTTCTTAAGCTG 416
Db 785 AKDTTTTTTTTWAATTTTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWT 844

QY 417 TGCAGTAATAGAGTATCGTCATTCATGTGATGATAAAGATGGAAGGGCTTCATTCAT 476
Db 845 WAATTTTATTAATATAWATAWTTTWWTTTAAWTTTWTWTWTWTWTWTWTWT 904

QY 477 GTTAGTATGGAATAGGAAGATAGGTAGGTAAGTATTTTAATAGATGTTTCTTTATGAAA 536
Db 905 WTNAAWNTAWAAWAAATTTATWTWTATYDWTWTATTTATTTTWTWTWTATWAKRA 964

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Qy 537 TAATTTTAAAGATTGTCAGCCCTGCATGATTTATGATCAATCATTTTGTGCTGCTT 596
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
965 WRWATWDAWAWRAATATTAATATATATATATATATATATATATATATATATATATAT 1024
Qy 597 AGTTACTTTTATAGAAATAGAAAGCATTTGTAGGCTCAGGAAAGCAAAACATTCAGAATGAA 656
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1025 AWDAATRTAAATWTWTTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTKTK 1084
Qy 657 ATCCAATAGAGAGGTA 673
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1085 TRTADATATATAATA 1101

RESULT 10
CNS016CO/c 1200 bp DNA linear GSS 26-JUL-1999
LOCUS Drosophila melanogaster genome survey sequence T7 end of BAC
DEFINITION BACN15E04 of DrosBAC library from Drosophila melanogaster (fruit
SOURCE fly), genomic survey sequence.
ACCESSION AL106578.1 GI:5622626.
KEYWORDS GSS.
ORGANISM Drosophila melanogaster (fruit fly)
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 1200)
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Barnot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Estimating of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
JOURNAL Nat. Genet. 25 (2), 235-238 (2000)
COMMENT Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk -. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaud at CEPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Payan. It has been constructed in the vector
pBelobAC11.
FEATURES             Location/Qualifiers
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                     /mol_type="genomic DNA"
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                     /plasmid="pBelobAC11"
                     /note="end : T7"

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Best Local Similarity 34.6%; Pred. No. 0.078;
Matches 121; Conservative 58; Mismatches 171; Indels 0; Gaps 0;

Qy 291 TAATAACCATGTCATCTAGAAATGAAAGTATATATTTTAAAGAGCATCGTTTGGCCATGT 350
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1175 TWTWATWAWWATATAATAATAATATWTTWAWWAAWAAATWATWATWATTTTANAATAT 1116
Qy 351 ATAAATTTTCAACACATTAACCTTCAGGGTTATTATTCCTTTTAGGCTAGTTTTCCTTA 410
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1115 WTWATWTTWTTWTTATTAATWTTTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWTWT 1056
Qy 411 AGTCGTGCAGTAATAGAGGTATCGTCATTCATGTCAGATAAAGATGGAAGGGGCTC 470
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1055 WTTWAWTWTWTTWAAATWATTTTATTTATTAATWAAATWTTTATWTTTATWTTATWTTW 996
Qy 471 ATTCAATGTAGTGGAATAGGAATAGGAATAGGTCAGTGAATGATTTTAAATAGATGTTCTTT 530
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
995 AAAAAAATWTTTWTWTTTAAATATAAARAWAATTAATTTWAAAWAATWTTTWTTRT 936
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Qy 531 ATGAATAATATTTTAAAGATTGTCAGCCCTGCATGATTTATGATCAATCATTTTGTGG 590
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935 TWWAATAATATTTTAAATATTTTCGTTAAWATATWTCHTTATTTTWTARAAATTTT 876
Qy 591 TCTGTTAGTTACTTTTATAGAAATAGAAAGCATTTGTAGGCTCAGGAAAGC 640
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
875 AATWTTTWTWTWTWTWTTATWTTWAAACCCDAAAAAAATTTWAGHAARGC 826

RESULT 11
CNS021G2 1146 bp DNA linear GSS 01-SEP-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 225004 of library G from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL176843
VERSION AL176843.1 GI:7814900
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontoidea; Tetraodontidae; Tetraodon.
REFERENCE 1
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Barnot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Estimating of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
JOURNAL Nat. Genet. 25 (2), 235-238 (2000)
MEDLINE 20296633
PUBMED 10835645
REFERENCE 2
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Ozouf-Costaz,C.,
Fizames,C., Bernot,A., Bouneau,L., Billault,A., Quetier,F.,
Saurin,W., Fierot,A. and Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Genome Res. 10 (7), 939-949 (2000)
MEDLINE 20359837
PUBMED 10899143
REFERENCE 3 (bases 1 to 1146)
AUTHORS Genoscope.
DIRECT SUBMISSION
SUBMITTED (12-APR-2000) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
FEATURES             Location/Qualifiers
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                     /mol_type="genomic DNA"
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                     /clone="225004"
                     /clone_lib="G"
                     /note="Genoscope sequence ID : COAG225BH02LP1-end : T7"

ORIGIN
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Best Local Similarity 25.6%; Pred. No. 0.097;
Matches 83; Conservative 106; Mismatches 131; Indels 4; Gaps 1;

Qy 452 AAAGATGGAAGGGCTTCATTCATGTTAGTATGGAATAAGAAATAGGAAATGATGAT 511
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
561 RAADAKRAAAKKTRWDTRATTTTADTAAAKTTAATTRRRKTTAAARRRRKRAKTRKK 620
Qy 512 TTTAATAGATGTTCTTTTATGAAATAATTTTAAAGATTCAGCCCTGCATGATTT 571
Db : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
621 TAKRRTTKDWAKTAARRKKTKRRKRRKTKTTTAAAGRKKTKTTTANKTAKTAATA 680
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978	ATAGTATTTTTTAAAAATTTATATATAGTAATTTATGCTTTTATGTTATTTTATATATTTTAT	1037
395	GGTCTAGTTTTTCTTAAAGTCGTGCGAGTAATAGAGGTATCGTCATTTCACTGTCACATAAAA	454
1038	TTTATATAGTTTTTATTTATTTTGTATATATATTTATTTTGTATATATATTTATTTTATATAG	1097
455	GATGGAAGGGGCTTCATTCATGCTTAGTCATAGGAAATAGGAAAGTAGGTGGAAGTGATTTT	514
1098	TTATTTTATTTTGTTTTATTTTATGGTATTTATTTATTTTGTATTTATTTATTTTAAATTT	1157
515	AATAGATGTTTCTTTTATGAAATPAAATTTTTTAAAGATTTGTCAGCCCTCGATGATTTATG	574
1158	TTTATATATTTATTTTATATATATAGTATTTATTTATATATTTATATATATGATTTAT	1217
575	ATGAATCATTTTGGTCTGTTAGTTACTTTTATAGAAAT	613
1218	TTTATAGATATTTTATATATTTATTTATTTATTTATTTATTTAT	1256

RESULT 13	AU088479/c	500 bp	linear	EST 27-JAN-2001
LOCUS	AU088479	Sugano Malaria cDNA library	Plasmodium falciparum	3D7 cDNA
DEFINITION	clone XFpN1713, mRNA sequence.			
VERSION	AU088479			
KEYWORDS	AU088479.1	GI:12390620		
SOURCE	EST.			
ORGANISM	Plasmodium falciparum	3D7		
REFERENCE	Plasmodium falciparum	3D7		
AUTHORS	Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.			
TITLE	1 (bases 1 to 500)			
JOURNAL	Watanabe,J., Sasaki,M., Suzuki,Y. and Sugano,S.			
MEDLINE	FULL-malaria: a database for a full-length enriched cDNA library			
PUBMED	from human malaria parasite, Plasmodium falciparum			
	Nucleic Acids Res. 29 (1), 70-71 (2001)			
	20574754			
	11125052			

Tel: 61-3-5449-5376
 Fax: 81-3-5449-5410
 Email: jwatanab@manage.ims.u-tokyo.ac.jp
 Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and
 Sugano, S. Construction and characterization of a full
 length-enriched and a 5'-end-enriched cDNA library Gene 200 (1-2),
 149-156 (1997)

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source
1. .500
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/mol_type="mRNA"
/isolate="3D7"
/db_xref="taxon:36329"
/clone="XPFn7173"
/dev stage="erythrocytic stage"
/clone_lib="Succano Malaria cdna library"

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Qy	396	GT	CT	AGT	CT	GT	CG	CA	GT	TA	T	AG	GT	AT	C	GT	CA	TT	CA	TT	CG	CA	TA	455
Db	360	TT	AT	AG	TT	TAT	AAAA	ATT	T	G	AA	AA	TT	T	A	A	G	T	T	A	T	A	T	301
Qy	456	AT	G	AA	A	G	G	G	CT	T	C	A	T	T	G	T	A	G	T	A	G	T	G	515


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Db      741 TATATATTTTTTTTTNTNTTATWAWWTTTTWTTTTTTTTTTATTTTATTTTATTTTATTTTATTTT 682
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Db      681 TATTTTTTATATTTTAWWTTTTWTTAWWWTTWTATWTTTWTATATAAATATATTTTATTTT 622
Qy      480 AGTGATGGAATAGGAAGTAGGTGAAGTGAATTTTAATAGATGTTCTTTTATGAAATAA 539
Db      621 ATWTTTATTTTGA--AAAAATATTTTTTTTTTAAAAATTTTTTATATTTATTTTAAATAAAAA 564
Qy      540 TTTTAAAGATTTGTCAGCCCTGCATGATTTATGATCAATTTTGTGGTCTGTTAGT 599
Db      563 ATATWAAAATWTTWTAWAATWTTTATTTATTTTAAAAAATTTTATTTTATTTTATTTTATTT 504
Qy      600 TACTTTTAGAATAGAAAAGCATTGTAGGCTCAGGGAAGCAAAACATTCAGAATGAAATC 659
Db      503 TWTWTTTAWAATWNTTTTAAATTTTATTAATTATAAAAAAAWAAWWTAAWRNTANTA 444
Qy      660 CAATAGA 666
Db      443 AAAAAA 437

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 Job time : 4017.41 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 8, 2005, 19:52:53 ; Search time 656.05 Seconds
(without alignments)
9232.381 Million cell updates/sec

Title: US-09-463-542-1_COPY_1_125
Perfect score: 125
Sequence: 1 cccctgcccctgccctgcc.....gcgggcagggggcccccagc 125

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues
Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl.*
1: gb.ba.*
2: gb.htg.*
3: gb.in.*
4: gb.on.*
5: gb.ov.*
6: gb.pat.*
7: gb.ph.*
8: gb.pl.*
9: gb.pr.*
10: gb.ro.*
11: gb.ats.*
12: gb.sy.*
13: gb.un.*
14: gb.vl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES				
Result No.	Score	Query Match %	Length	ID Description
1	125	100.0	308	9 AF012873 Homo sapi
2	118.6	94.9	135675	9 AC093174 Homo sapi
3	118.6	94.9	180963	9 AC091492 Homo sapi
4	117	93.6	148828	9 AV157024 Homo sapi
5	62.2	49.8	108399	2 AL356736 Homo sapi
6	61	48.8	238794	2 BX936377 Homo sapi
7	59.6	47.7	171574	2 AC012300 Homo sapi
8	59.4	47.5	135119	2 AC011578 Homo sapi
9	58.6	46.9	53121	2 AC136331 Homo sapi
10	58.6	46.9	68330	2 AC116109 Mus muscu
11	58.2	46.6	72645	2 AC112672 Mus muscu
12	58.2	46.6	80542	2 AC020683 Homo sapi
13	58	46.4	84701	2 AC016122 Homo sapi
14	57.8	46.2	1416	6 AX840138 Sequence
15	57.8	46.2	1416	6 AX840139 Sequence
16	57.8	46.2	234009	2 CR383672 Danio rer
17	57.6	46.1	75449	2 AC016188 Homo sapi
18	57.6	46.1	79023	2 AC021951 Homo sapi
19	57.4	45.9	59727	2 AC100218 Mus muscu

c	20	57.4	45.9	212028	2	CR751564	CR751564
	21	57.4	45.9	218236	2	AC117823	Danio rer
	22	57.4	45.9	240965	2	AC126054	Mus muscu
	23	57.2	45.8	87440	2	AC021226	AC126054 Mus muscu
	24	57.2	45.8	218311	2	CR354540	AC021226 Homo sapi
	25	56.8	45.4	1060	11	BV111230	CR354540 Danio rer
c	26	56.8	45.4	126667	2	AC015839	BV111230 PZA00983
	27	56.6	45.3	66680	2	AC100483	AC015839 Homo sapi
	28	56.6	45.3	72847	2	AC023563	AC100483 Mus muscu
	29	56.6	45.3	87283	2	AC022856	AC023563 Homo sapi
c	30	56.6	45.3	180652	2	CR762436	AC022856 Homo sapi
	31	56.4	45.1	78700	2	AC022685	CR762436 Danio rer
	32	56.4	45.1	110000	2	AL929091_06	AC022685 Homo sapi
c	33	56.4	45.1	130244	2	AC013518	Continuation (7 of
c	34	56.4	45.1	186731	2	AC073212	AC013518 Homo sapi
	35	56.4	45.1	194034	9	AL592148	AC073212 Homo sapi
	36	56.2	45.0	1703	9	HSHB2A	AL592148 Human DNA
	37	56.2	45.0	2904	9	HSHB2G	X74143 H.sapiens H
	38	56.2	45.0	82815	2	AC062001	X78202 H.sapiens H
	39	56	44.8	51274	2	AC023454	AC062001 Homo sapi
	40	56	44.8	229518	2	AC069058	AC023454 Homo sapi
c	41	55.8	44.6	3281	2	AC143602	AC069058 Homo sapi
c	42	55.8	44.6	66596	2	AC131276	AC143602 Macaca mu
c	43	55.8	44.6	78220	2	AC023212	AC131276 Homo sapi
	44	55.8	44.6	81767	2	AC021929	AC023212 Homo sapi
	45	55.8	44.6	132745	2	AC134836	AC021929 Homo sapi
							AC134836 Mus muscu

ALIGNMENTS

RESULT 1
AF012873
LOCUS AF012873 308 bp DNA linear PRI 03-MAY-1999
DEFINITION Homo sapiens PPARG gene, isoform 1, promoter and 5'UTR sequence.
ACCESSION AF012873
VERSION AF012873.1 GI:4731427
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 308)
AUTHORS Fajas,L., Auboeuf,D., Raspe,E., Schoonjans,K., Lefebvre,A.-M.,
Saladin,R., Najib,J., Laville,M., Fruchart,J.-C., Deeb,S.,
Puig-Vidal,A., Flier,J., Briggs,M., Vidal,H., Staels,B. and
Auwerx,J.
TITLE The organization, promoter analysis, and expression of the human
PPARG gene
JOURNAL J. Biol. Chem. 272 (30), 18779-18789 (1997)
MEDLINE 97373577
PUBMED 9228052
REFERENCE 2 (bases 1 to 308)
AUTHORS Fajas,L., Auwerx,J., Saladin,R. and Briggs,M.
TITLE Direct Submission
JOURNAL Submitted (07-JUL-1997) Cardiovascular Res., Ligand
Pharmaceuticals, 9393 Towne Centre Dr., San Diego, CA 92121-3016,
USA
FEATURES
source Location/Qualifiers
1..308
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
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1..126
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127..308
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promoter
5'UTR
ORIGIN


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Db          20832 CC 20833
RESULT 8
AC011578/c
LOCUS       AC011578      135119 bp      DNA      linear      HTG 13-JUL-2000
DEFINITION Homo sapiens clone RP11-12P9, LOW-PASS SEQUENCE SAMPLING.
ACCESSION   AC011578
VERSION     AC011578.4 GI:9123847
KEYWORDS    HTG; HTGS PHASE0.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 135119)
AUTHORS     Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE       Homo sapiens, clone RP11-12P9
JOURNAL
REFERENCE   2 (bases 1 to 135119)
AUTHORS     Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
            Baldwin,J., Barna,N., Beckerly,R., Boguslavskiy,L., Bouckghalter,B.,
            Brown,A., Castle,A., Collangelo,M., Collins,S., Collymore,A.,
            Cooke,P., Dearellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,
            Ferreira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D., Horton,L.,
            Galagan,J., Gardyna,S., Grant,G., Hegos,B., Heaford,A., Klein,J.,
            Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
            Lehocsky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
            McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
            Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
            Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
            Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
            Testaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
            Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
            Direct Submission
            Submitted (07-OCT-1999) Whitehead Institute/MIT Center for Genome
            Research, 320 Charles Street, Cambridge, MA 02141, USA
            On Jul 13, 2000 this sequence version replaced gi:7107923.
            All repeats were identified using RepeatMasker:
            Smit, A.F.A. & Green, P. (1996-1997)
            http://ftp.genome.washington.edu/RM/RepeatMasker.html
            ----- Genome Center
            Center: Whitehead Institute/ MIT Center for Genome Research
            Center code: WIGR
            Web site: http://www-seq.wi.mit.edu
            Contact: sequence_submissions@genome.wi.mit.edu
            ----- Project Information
            Center project name: L3313
            Center clone name: 12_P_9
            -----
            * NOTE: This record contains 146 individual
            * sequencing reads that have not been assembled into
            * contigs. Runs of N are used to separate the reads
            * and the order in which they appear is completely
            * arbitrary. Low-pass sequence sampling is useful for
            * identifying clones that may be gene-rich and allows
            * overlap relationships among clones to be deduced.
            * However, it should not be assumed that this clone
            * will be sequenced to completion. In the event that
            * the record is updated, the accession number will
            * be preserved.
            *
            * 1 744: contig of 744 bp in length
            * 745 844: gap of 100 bp
            * 845 1615: contig of 771 bp in length
            * 1616 1715: gap of 100 bp
            * 1716 2485: contig of 770 bp in length
            * 2486 2585: gap of 100 bp
            * 2586 3345: contig of 760 bp in length
            * 3346 3445: gap of 100 bp
            * 3446 4188: contig of 743 bp in length
            * 4189 4288: gap of 100 bp
            * 4289 5050: contig of 762 bp in length
            * 5051 5150: gap of 100 bp
            * 5151 5984: contig of 834 bp in length
            *
            * 5985 6084: gap of 100 bp
            * 6085 6837: contig of 753 bp in length
            * 6838 7694: contig of 757 bp in length
            * 7695 7794: gap of 100 bp
            * 7795 8541: contig of 747 bp in length
            * 8542 8641: gap of 100 bp
            * 8642 9403: contig of 762 bp in length
            * 9404 9503: gap of 100 bp
            * 9504 10273: contig of 770 bp in length
            * 10274 10373: gap of 100 bp
            * 10374 11138: contig of 765 bp in length
            * 11139 11238: gap of 100 bp
            * 11239 11991: contig of 753 bp in length
            * 11992 12091: gap of 100 bp
            * 12092 12840: contig of 749 bp in length
            * 12841 12940: gap of 100 bp
            * 12941 13800: contig of 860 bp in length
            * 13801 13901: gap of 100 bp
            * 13901 14661: contig of 761 bp in length
            * 14662 14761: gap of 100 bp
            * 14762 15508: contig of 747 bp in length
            * 15509 15608: gap of 100 bp
            * 15609 16360: contig of 751 bp in length
            * 16360 16460: gap of 100 bp
            * 16460 17225: contig of 766 bp in length
            * 17226 17325: gap of 100 bp
            * 17326 18074: contig of 749 bp in length
            * 18075 18174: gap of 100 bp
            * 18175 18916: contig of 742 bp in length
            * 18917 19016: gap of 100 bp
            * 19017 19785: contig of 769 bp in length
            * 19786 19886: gap of 100 bp
            * 19886 20641: contig of 756 bp in length
            * 20642 20741: gap of 100 bp
            * 20742 21501: contig of 760 bp in length
            * 21502 21601: gap of 100 bp
            * 21602 22367: contig of 766 bp in length
            * 22368 22467: gap of 100 bp
            * 22468 23213: contig of 746 bp in length
            * 23214 23313: gap of 100 bp
            * 23314 24069: contig of 756 bp in length
            * 24070 24169: gap of 100 bp
            * 24170 24922: contig of 753 bp in length
            * 24923 25022: gap of 100 bp
            * 25023 25771: contig of 749 bp in length
            * 25772 25871: gap of 100 bp
            * 25872 2632: contig of 761 bp in length
            * 26633 27332: gap of 100 bp
            * 27333 27495: contig of 763 bp in length
            * 27496 27595: gap of 100 bp
            * 27596 28366: contig of 771 bp in length
            * 28367 28466: gap of 100 bp
            * 28467 29222: contig of 756 bp in length
            * 29223 29323: gap of 100 bp
            * 29323 30092: contig of 770 bp in length
            * 30093 30192: gap of 100 bp
            * 30193 30945: contig of 753 bp in length
            * 30946 31045: gap of 100 bp
            * 31046 31793: contig of 748 bp in length
            * 31794 31893: gap of 100 bp
            * 31894 32649: contig of 756 bp in length
            * 32650 32749: gap of 100 bp
            * 32750 33490: contig of 741 bp in length
            * 33491 33590: gap of 100 bp
            * 33591 34331: contig of 741 bp in length
            * 34332 34431: gap of 100 bp
            * 34432 35176: contig of 745 bp in length
            * 35177 35276: gap of 100 bp
            * 35277 36039: contig of 763 bp in length
            * 36040 36139: gap of 100 bp
            * 36140 36886: contig of 747 bp in length
            * 36887 36986: gap of 100 bp

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RESULT 10
AC116109
LOCUS      AC116109               68330 bp    DNA        linear    HTG 25-MAR-2002
DEFINITION Mus musculus clone RP23-467G14, LOW-PASS SEQUENCE SAMPLING.
ACCESSION  AC116109
VERSION    AC116109.1  GI:19703220
KEYWORDS   HTG; HTGS PHASE0.
SOURCE     Mus musculus (house mouse)
ORGANISM   Mus musculus

REFERENCE  1  (bases 1 to 68330)
AUTHORS   Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE     Mus musculus, clone RP23-467G14
JOURNAL   Unpublished
REFERENCE  2  (bases 1 to 68330)
AUTHORS   Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
          Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L.,
          Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
          Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
          Cook,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S., Dodge,S.,
          Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
          Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
          Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
          Kamat,A., Karatas,A., Kelle,C., LaRocque,K., Lamazares,R.,
          Landers,T., Lehoczeky,J., Levine,R., Lindblad-Toh,K., Liu,G.,
          MacLean,C., MacDonald,P., Major,O., Marquis,N., Matthews,C.,
          McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Meneus,L.,
          Mihovaty,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R.,
          Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
          Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
          Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
          Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupack,R.,
          Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
          Strausman,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
          Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
          Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
          Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
TITLE     Direct Submission
JOURNAL   Submitted (25-MAR-2002) Whitehead Institute/MIT Center for Genome
          Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT   All repeats were identified using RepeatMasker:
          Smit, A.F.A. & Green, P. (1996-1997)
          http://ftp.genome.washington.edu/RM/RepeatMasker.html
          -----
          Center: Genome Center
          Center: Whitehead Institute/ MIT Center for Genome Research
          Center code: WIBR
          Web site: http://www-seq.wi.mit.edu
          Contact: sequence_submissions@genome.wi.mit.edu
          -----
          Project Information
          Center project name: L22810
          Center clone name: 467_G_14
          -----
          * NOTE: This record contains 85 individual
          * sequencing reads that have not been assembled into
          * contigs. Runs of N are used to separate the reads
          * and the order in which they appear is completely
          * arbitrary. Low-pass sequence sampling is useful for
          * identifying clones that may be gene-rich and allows
          * overlap relationships among clones to be deduced.
          * However, it should not be assumed that this clone
          * will be sequenced to completion. In the event that
          * the record is updated, the accession number will
          * be preserved.
          * 1
          * 648 747: contig of 647 bp in length
          * 748 1422: contig of 675 bp in length
          * 1423 1522: gap of 100 bp
          * 1523 2229: contig of 707 bp in length
          * 2230 2329: gap of 100 bp
          * 2330 3001: contig of 672 bp in length
          * 3002 3101: gap of 100 bp
          *
          * 3102 contig of 682 bp in length
          * 3784 gap of 100 bp
          * 3884 contig of 703 bp in length
          * 4587 gap of 100 bp
          * 4687 contig of 695 bp in length
          * 5381 gap of 100 bp
          * 5382 contig of 709 bp in length
          * 6190 gap of 100 bp
          * 6191 contig of 713 bp in length
          * 6291 gap of 100 bp
          * 7004 gap of 100 bp
          * 7104 contig of 721 bp in length
          * 7825 gap of 100 bp
          * 7925 contig of 738 bp in length
          * 8663 gap of 100 bp
          * 8763 contig of 640 bp in length
          * 9403 gap of 100 bp
          * 9503 contig of 733 bp in length
          * 10236 gap of 100 bp
          * 10336 contig of 720 bp in length
          * 11056 gap of 100 bp
          * 11156 contig of 706 bp in length
          * 11861 gap of 100 bp
          * 11862 contig of 692 bp in length
          * 12653 gap of 100 bp
          * 12754 contig of 692 bp in length
          * 13446 gap of 100 bp
          * 13446 contig of 690 bp in length
          * 14236 gap of 100 bp
          * 14336 contig of 711 bp in length
          * 15047 gap of 100 bp
          * 15147 contig of 707 bp in length
          * 15853 gap of 100 bp
          * 15953 contig of 679 bp in length
          * 16632 gap of 100 bp
          * 16732 contig of 709 bp in length
          * 17441 gap of 100 bp
          * 17541 contig of 729 bp in length
          * 18270 gap of 100 bp
          * 18370 contig of 678 bp in length
          * 19048 gap of 100 bp
          * 19148 contig of 703 bp in length
          * 19449 gap of 100 bp
          * 19851 contig of 710 bp in length
          * 20661 gap of 100 bp
          * 20662 contig of 712 bp in length
          * 21473 gap of 100 bp
          * 21474 contig of 713 bp in length
          * 21574 gap of 100 bp
          * 22287 contig of 700 bp in length
          * 22387 gap of 100 bp
          * 23087 contig of 734 bp in length
          * 23187 gap of 100 bp
          * 23921 contig of 693 bp in length
          * 24021 gap of 100 bp
          * 24714 contig of 727 bp in length
          * 24814 gap of 100 bp
          * 25541 contig of 724 bp in length
          * 25641 gap of 100 bp
          * 26365 contig of 716 bp in length
          * 26465 gap of 100 bp
          * 27181 contig of 697 bp in length
          * 27281 gap of 100 bp
          * 27978 contig of 692 bp in length
          * 28078 gap of 100 bp
          * 28770 contig of 690 bp in length
          * 28870 gap of 100 bp
          * 29560 contig of 698 bp in length
          * 29660 gap of 100 bp
          * 30358 contig of 708 bp in length
          * 30458 gap of 100 bp
          * 31166 contig of 683 bp in length
          * 31266 gap of 100 bp
          * 31949 contig of 720 bp in length
          * 32049 gap of 100 bp

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Wed Nov 9 09:34:50 2005

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AC020683      AC020683      80542 bp      DNA      linear      HTG 13-JUL-2000
LOCUS         Homo sapiens clone RP11-30L16, LOW-PASS SEQUENCE SAMPLING.
DEFINITION
AC020683
AC020683      AC020683.2  GI:9140295
VERSION       HTG; HTGS PHASE0
KEYWORDS
SOURCE        Homo sapiens (human)
ORGANISM      Homo sapiens
REFERENCE     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS       Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE        1 (bases 1 to 80542)
JOURNAL
REFERENCE     Birren, B., Linton, L., Nusbaum, C. and Lander, E.
AUTHORS       2 (bases 1 to 80542)
        Birren, B., Linton, L., Nusbaum, C., Lander, E., Abraham, H., Allen, N.,
        Anderson, S., Baldwin, J., Barna, N., Beckerly, R., Bada, F.,
        Boguslavskiy, L., Boukhgalter, B., Brown, A., Burkett, G., Castie, A.,
        Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P.,
        Dearellano, K., Dewar, K., Domino, M., Doyle, M., Fenestor, J.,
        Ferreira, P., FitzHugh, W., Forrest, C., Gage, D., Galagan, J.,
        Gardyna, S., Grant, G., Hagos, B., Heaford, A., Horton, L.,
        Howland, J. C., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J.,
        Landers, T., Lehoczy, J., Levine, R., Lieu, C., Liu, G., Locke, K.,
        MacDonald, P., Marquis, N., McEwan, P., McGurk, A., McKernan, K.,
        McPheeters, R., Meldrim, J., Meneus, L., Morrow, J., Naylor, J.,
        Norman, C. H., O'Connor, T., O'Donnell, P., Olivari, T. M., Peterson, K.,
        Pierre, N., Pisani, C., Pollara, V., Raymond, C., Riley, R., Rothman, D.,
        Roy, A., Santos, R., Severy, P., Spencer, B., Stange-Thomann, N.,
        Stojanovic, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
        Tirrell, A., Vasiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W. J.,
        Zimmer, A. and Zody, M.
        Direct Submission
        Submitted (08-JAN-2000) Whitehead Institute/MIT Center for Genome
        Research, 320 Charles Street, Cambridge, MA 02141, USA
        On Jul 13, 2000 this sequence version replaced gi:6682660.
        All repeats were identified using RepeatMasker:
        Smit, A.F.A. & Green, P. (1996-1997)
        http://ftp.genome.washington.edu/RM/RepeatMasker.html
        ----- Genome Center
        Center: Whitehead Institute/ MIT Center for Genome Research
        Center code: WIBR
        Web site: http://www-seq.wi.mit.edu
        Contact: sequence_submissions@genome.wi.mit.edu
        ----- Project Information
        Center project name: L4895
        Center clone name: 30_L_16
        -----
        * NOTE: This record contains 93 individual
        * sequencing reads that have not been assembled into
        * contigs. Runs of N are used to separate the reads
        * and the order in which they appear is completely
        * arbitrary. Low-pass sequence sampling is useful for
        * identifying clones that may be gene-rich and allows
        * overlap relationships among clones to be deduced.
        * However, it should not be assumed that this clone
        * will be sequenced to completion. In the event that
        * the record is updated, the accession number will
        * be preserved.
        *
        * 1      751: contig of 751 bp in length
        * 752      851: gap of 100 bp
        * 852      1603: contig of 752 bp in length
        * 1604      1703: gap of 100 bp
        * 1704      2545: contig of 842 bp in length
        * 2546      2645: gap of 100 bp
        * 2646      3412: contig of 767 bp in length
        * 3413      3512: gap of 100 bp
        * 3513      4285: contig of 773 bp in length
        * 4286      4386: gap of 100 bp
        * 4386      5154: contig of 769 bp in length
        * 5155      5254: gap of 100 bp
        * 5255      6018: contig of 764 bp in length
        * 6019      6118: gap of 100 bp
        *
        * 6119      6884: contig of 766 bp in length
        * 6885      6984: gap of 100 bp
        * 6985      7755: contig of 771 bp in length
        * 7756      7855: gap of 100 bp
        * 7856      8625: contig of 770 bp in length
        * 8626      8725: gap of 100 bp
        * 8726      9469: contig of 744 bp in length
        * 9470      9569: gap of 100 bp
        * 9570      10338: contig of 769 bp in length
        * 10339      10438: gap of 100 bp
        * 10439      11225: contig of 787 bp in length
        * 11226      11325: gap of 100 bp
        * 11326      12098: contig of 773 bp in length
        * 12099      12198: gap of 100 bp
        * 12199      12973: contig of 775 bp in length
        * 12974      13073: gap of 100 bp
        * 13074      13831: contig of 758 bp in length
        * 13832      13931: gap of 100 bp
        * 13932      14698: contig of 767 bp in length
        * 14699      14798: gap of 100 bp
        * 14799      15569: contig of 771 bp in length
        * 15570      15669: gap of 100 bp
        * 15670      16447: contig of 778 bp in length
        * 16448      16547: gap of 100 bp
        * 16548      17303: contig of 756 bp in length
        * 17304      17403: gap of 100 bp
        * 17404      18172: contig of 769 bp in length
        * 18173      18272: gap of 100 bp
        * 18273      19043: contig of 771 bp in length
        * 19044      19143: gap of 100 bp
        * 19144      19900: contig of 757 bp in length
        * 19901      20000: gap of 100 bp
        * 20001      20769: contig of 769 bp in length
        * 20770      20869: gap of 100 bp
        * 20870      21648: contig of 779 bp in length
        * 21649      21748: gap of 100 bp
        * 21749      22522: contig of 774 bp in length
        * 22523      22622: gap of 100 bp
        * 22623      23389: contig of 767 bp in length
        * 23390      23489: gap of 100 bp
        * 23490      24262: contig of 773 bp in length
        * 24263      24362: gap of 100 bp
        * 24363      25128: contig of 766 bp in length
        * 25129      25228: gap of 100 bp
        * 25229      25989: contig of 761 bp in length
        * 25990      26089: gap of 100 bp
        * 26090      26861: contig of 772 bp in length
        * 26862      26961: gap of 100 bp
        * 26962      27737: contig of 776 bp in length
        * 27738      27837: gap of 100 bp
        * 27838      28612: contig of 775 bp in length
        * 28613      28712: gap of 100 bp
        * 28713      29477: contig of 765 bp in length
        * 29478      29577: gap of 100 bp
        * 29578      30327: contig of 750 bp in length
        * 30328      30427: gap of 100 bp
        * 30428      31195: contig of 768 bp in length
        * 31196      31295: gap of 100 bp
        * 31296      32066: contig of 771 bp in length
        * 32067      32166: gap of 100 bp
        * 32167      32942: contig of 776 bp in length
        * 32943      33042: gap of 100 bp
        * 33043      33818: contig of 776 bp in length
        * 33819      33918: gap of 100 bp
        * 33919      34680: contig of 762 bp in length
        * 34681      34780: gap of 100 bp
        * 34781      35554: contig of 774 bp in length
        * 35555      35654: gap of 100 bp
        * 35655      36512: contig of 758 bp in length
        * 36513      36413      37292: contig of 780 bp in length
        * 37293      37392: gap of 100 bp
        * 37393      38153: contig of 761 bp in length

```


